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Development of novel LL-37 derived antimicrobial peptides with LPS and LTA neutralizing and antimicrobial activities for therapeutic application

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1. Introduction

Chronic upper airway infections affect a large part of the population [45]. Children may suffer from long-term effects with respect to the development of speech and cognitive abilities because of the hearing loss that accompanies otitis media with effusion (OME) [16]. Chronic sinusitis seriously affects the state of well being of these patients [6,14]. Nowadays, most patients with upper airway infections are treated with antibiotics. However, due to the growing resistance of bacteria for conventional antibiotics [17] it is necessary to develop alternative and more efficient strategies in the battle against chronic upper airway infections.

In a large part of these infections bacterial products, like lipopolysaccharide (LPS) from Gram-negative bacteria and lipoteichoic acid (LTA) from Gram-positive bacteria, are involved [15,28]. These products can induce an inflammatory reaction in the middle ear or in the sinuses and can cause injury to the mucosa of the upper airway epithelia [25,26].

Antimicrobial peptides play an important role in innate host defense, and this is believed to be particularly important at mucosal surfaces that form the initial barrier between the
Defensins and cathelicidins are the principal human antimicrobial peptides. They are mainly produced by neutrophils and epithelial cells, and have the capacity to kill and/or inactivate bacteria, fungi, and enveloped viruses in vitro [18,47]. The only member of the cathelicidin family identified in humans is hCAP18 (human cationic antibacterial protein of 18 kDa), and its carboxy-terminal antibacterial peptide, called LL-37, which comprises 37 amino acid residues [12]. hCAP18 is present in neutrophils, and is expressed throughout epithelia in many organs, including surface epithelia of the airways and submucosal glands [2]. LL-37 is considered to play an important role in the first line of defense against local and systemic infection and in systemic invasion of pathogens at sites of inflammation and wounds [3]. It has been shown to have a broad antimicrobial activity and it is able to neutralize LPS and LTA [34,42]. On the other hand, LL-37 has also been shown to have chemotactic effects on host cells [9,27].

Since LL-37 is suggested to play a role in innate host defense at mucosal surfaces, we investigated the therapeutic potential of synthetic peptides that we developed based on the sequence of LL-37. It has been suggested that the activity of LL-37 is related to its amphipatic helical structure. We therefore developed length variants with predicted improved amphipatic character.

Our aim was to obtain shorter peptides with improved or similar LPS and LTA neutralizing activities as LL-37, but with lower chemotactic properties. The efficacy was measured by LPS neutralization in a limulus amoebocyte lysate (LAL) assay and a whole blood IL-8 release assay (WB assay). LTA neutralization was also measured with a WB assay. To determine whether our synthetic peptides, designed to neutralize inflammation, would potentially promote immunity and inflammation we assessed their capacity to activate T cells (in an Elispot and a T cell proliferation assay), tested extracellular signal-regulated kinase (ERK)-activation of epithelial cells, and measured chemotaxis of neutrophils. In addition, we studied the toxicity and the antibacterial and antifungal activity of a selected peptide.

2. Materials and methods

2.1. Preparation and characterization of synthetic peptides

2.1.1. Peptide development

To develop peptide derivatives of LL-37 with improved or similar LPS and LTA neutralizing activities, we first analyzed 20 different 22-amino acid peptides spanning the LL-37 sequence (Table 1). The peptides were tested for their LPS neutralizing activity in a limulus amoebocyte lysate (LAL) assay and a whole blood IL-8 release assay (WB assay). LTA neutralization was also measured with a WB assay. To determine whether our synthetic peptides, designed to neutralize inflammation, would potentially promote immunity and inflammation we assessed their capacity to activate T cells (in an Elispot and a T cell proliferation assay), tested extracellular signal-regulated kinase (ERK)-activation of epithelial cells, and measured chemotaxis of neutrophils. In addition, we studied the toxicity and the antibacterial and antifungal activity of a selected peptide.
and LTA neutralizing capacities. From the four most potent peptides we synthesized one 25-mer, two 24-mers, and three 23-mers. One of the 24-mers showed good LPS and LTA neutralizing properties and was called P60. From this peptide, 10 variants were made with several amino acid modifications in order to increase amphipaticity and stability. In Fig. 1 helical wheel representations of P60 and P60.4 are shown to illustrate the expected enhanced amphipaticity. Variant P60.4 and P60 showed the best neutralizing capacities and were further investigated in detail in this study. A more stable variant of P60, wherein the N-terminal part is acetylated and the C-terminal part is amidated (P60-Ac) was also studied. For the toxicity studies, P60.4 in acetylated and amidated form (P60.4-Ac) was used. In the otoxicity study the peptide was solved in a formulation solution (7% Macrogol 10,000 in isotope [NaCl] preserved [0.02% benzalkonium chloride and 0.1% Na2EDTA] 20 mM phosphate pH 5.5), which was developed to serve as future eardrop formulation.

2.1.2. Peptide synthesis
Peptides were synthesized by solid phase strategies on an automated multiple peptide synthesizer (Syroff, MultiSynTech, Witten, Germany). Tentagel S AC (Rapp, Tübingen, Germany), a graft polymer of polyethylene glycol and polystyrene was used as a resin (loading 0.2 meq, particle size 90 μm) [31,35]. Repetitive couplings were performed by adding a 6-fold molar excess (based on the resin loading) of a 0.60 M solution of the appropriate 9-fluorenylmethoxycarbonyl (Fmoc) amino acid [7,10,11] in N-methylpyrrolidone (NMP), a 6-fold molar excess of 0.67 M benzotriazole-1-yl-oxy-tris-pyrrolidino-phosphonium hexafluorophosphate (PyBOP) [8] in NMP and a 12-fold molar excess of N-methylmorpholine (NMM) in NMP 2/1 (v/v) to each reaction vessel. Coupling- and deprotection-times were 45 min and three times 3 min, respectively. Washings after couplings and Fmoc-deprotection were performed six times with NMP. Amidation of the peptide was obtained by using Tantagel S AM (Rapp, Tübingen, Germany), acetylation was performed by coupling acetic acid. After synthesis the peptidyl resins were washed extensively with NMP, dichloromethane, dichloromethane/ether 1/1 (v/v) and ether, respectively, and air-dried. Peptidyl resins were cleaved and side chain deprotected in trifluoroacetic acid (TFA)/water 95/5 (v/v) for 2.5 h (1.5 ml/10 μmol peptide), the resin was removed by filtration and the peptide was precipitated from the TFA solution with ether/pentane 1/1 (v/v) (10 ml/10 μmol peptide). The solution was cooled for 1 h at −20 °C and the precipitated peptide was isolated by centrifugation (−20 °C, 2500 × g, 10 min.). After washing and vortexing of the pellet with 10 ml ether/pentane 1/1 (v/v) and isolation by the same procedure, the peptides were air dried at room temperature for 1 h. Peptides were dissolved in 2 ml water or 2 ml 10 vol% acetic acid, the solution was frozen in liquid nitrogen for about 5 min and subsequently lyophilized while being centrifuged (1300 rpm, 8–16 h).

2.1.3. RP-HPLC analysis
Lyophilized peptides were dissolved in 10 vol% acetic acid (1 mg/ml) and a sample of 20–50 μl was analyzed on a Waters LCM-1 system equipped with a CC 125/4 nucleosil 100-5 C18 AB column (Macherey-Nagel, Düren, Germany). A ternary gradient system was used of A: water, B: acetonitrile, and C: 2 vol% TFA in water. For RP-HPLC analysis a linear gradient was applied of A/B/C 90/5/5 to A/B/C 20/75/5 in 22 min. The flow was 1.0 ml/min and detection was at 214 nm. Peak integration of the chromatogram between 4 and 22 min was performed and peptide purity was defined as the peak area of the peptide compared to the total peak area.

2.1.4. Maldi-Tof mass spectrometry analysis
The peptide solutions were diluted 1:100 with acetonitrile/water/TFA 50/50/0.2 (v/v/v) and 1 μl of each solution was mixed with 1 μl of a 10 mg/ml solution of α-cyano-4-hydroxycinnamic acid (4-ACH) in acetonitrile/water/TFA 50/50/0.2 (v/v/v) containing a mass reference peptide (VNTPEHVPYGLGSPSRS, bovine big-endothelin (22–39), MH_u* = 1897.10). The mixtures were applied to a sample slide, allowed to dry and crystallize (about 10 min) and applied to the mass spectrometer (Voyager DE-Pro, PerSeptive Biosystems, Framingham, USA). The instrument was used in linear mode with an acquisition mass range of 500–5000 Da and 50 shots/spectrum (3 Hz) were applied at a laser intensity of 2000. Measured average masses were corrected based on the mass of the reference peptide and subsequently compared to calculated average masses.

2.2. LPS neutralization assays
The synthetic peptides were examined for LPS neutralizing activity in a limulus amoebocyte lysate (LAL) assay and in a...
whole blood assay. For the LAL assay, peptides in concentrations from 0.125 to 16 μg/ml, were incubated with 1 ng/ml LPS (from Salmonella typhi; Sigma Chemicals, Zwijndrecht, The Netherlands) for 30 min at 37 °C in a 96-well plate (#3596; Costar, Cambridge, MA). Residual LPS activity was determined using the LAL assay (Bio Whittaker Inc., Walkersville, MD).

For the whole blood assay, heparinized blood was collected by venipuncture into heparinized tubes (Becton Dickinson, Plymouth, UK). After preincubation of 25 μl 500 ng/ml LPS (final concentration 1 ng/ml) with 25 μl peptide (final concentrations 0.125–16 μg/ml) for 30 min at 37 °C, blood (200 μl) was added in a 96-well plate (#655180; Greiner, Alphen a/d Rijn, The Netherlands). Subsequently, the blood was stimulated and incubated with the LPS mixture for 24 h at 37 °C. After centrifugation for 8 min at 1200 rpm, plasma was collected and stored at −20 °C until analysis for IL-8 levels by ELISA (BioSource Europe S.A., Nivelles, Belgium).

2.3. LTA neutralization assays

For the LTA neutralization, a whole blood assay was performed as described for LPS neutralization. Peptides were preincubated in a final concentration of 50 ng/ml LTA (from Staphylococcus aureus; Sigma Chemicals, Zwijndrecht, The Netherlands). Plasma was stored at −20 °C until analysis for IL-8 levels by ELISA (BioSource Europe S.A., Nivelles, Belgium).

2.4. T cell assays

2.4.1. Isolation of PBMC from heparinized blood

To test the synthetic peptides in T cell assays, heparinized blood was acquired from three different families, which included a total of four adults and four children. Peripheral blood mononuclear cells (PBMC) were obtained by Ficoll gradient centrifugation by layering 10 ml of Ficoll under 35 ml of PBS-diluted heparinized blood. PBMC were collected from the interface and washed twice with PBS before dilution in Iscove’s Modified Dulbecco’s Medium (Gibco Life Technologies, Breda, The Netherlands) containing 10% pooled human AB serum, 2 mM glutamine (Bio Whittaker Inc., Walkersville, MD), 100 U/ml penicillin (Bio Whittaker), 20 μg/ml streptomycin (Bio Whittaker Inc.) and 100 IU/ml protease inhibitor cocktail (Boeringer Mannheim, Roche, Basel, Switzerland). Samples were subjected to SDS-PAGE on a 10% glycin-based gel. Resolved proteins were transferred to a polyvinylidene difluoride (PVDF) membrane, and non-specific binding sites were blocked by PBS/0.05% Tween-20/1% casein. The blots were incubated with rabbit polyclonal antibodies against phosphorylated ERK-1/2 (New England Biolabs, Beverly, MA), and secondary horseradish peroxidase conjugated anti-rabbit IgG antibodies. The enhanced chemoluminescent (ECL) Western blotting detection system (Amersham Pharmacia Biotech, Uppsala, Sweden) was used to reveal immunoreactivity.

2.4.2. T cell proliferation assay

For T cell proliferation, 150,000 PBMC were cultured in the absence or presence of various concentrations (1, 10, and 100 μg/ml) of synthetic peptides for 5 days in 96-well round bottom plates (Costar Inc. Cambridge, MA) in a final volume of 150 μl IMDM complete. As a positive control, PBMC were cultured in the presence of 25 U/ml recombinant IL-2. During the final 20 h of culture, PBMC were pulsed with [3H] thymidine (0.5 μCi/well), after which 3H-incorporation was measured by liquid scintillation counting. Results are expressed as mean counts per minute (cpm) of triplicate wells.

2.4.3. Elistop assay

For detection of the T cell cytokines IFN-γ and IL-10 by Elistop analysis, 1.5 × 10^6 PBMC were cultured in 0.5 ml IMDM complete in the absence or presence of various concentrations of synthetic peptide (2.5 and 25 μg/ml). As a positive control PBMC were stimulated with 10 μg/ml pokeweed mitogen (PWM). After 48 h of culture, PBMC were harvested by gently rinsing the wells with warm IMDM to collect non-adherent cells, which were washed in a large volume of IMDM. PBMC were subsequently plated on antibody-precoated ELISA plates and cultured for 5 h in IMDM supplemented with 2% pooled human AB serum at 37 °C 5% CO2, after which the plates were developed according to the manufacturer’s protocol (U-GyTech, Utrecht, The Netherlands). Results are expressed as mean spot numbers per well of triplicate wells.

2.5. ERK1/2-activation

Cells from the muco-epidermoid lung carcinoma cell line NCI-H292 (ATCC, Rockville, MD) were cultured in 24- or 6-well tissue culture plates in RPMI1640 medium (Gibco, Grand Island, NY) supplemented with 2 mM l-glutamine (Bio Whittaker, Walkersville, MD), 20 U/ml penicillin (Bio Whittaker), 20 μg/ml streptomycin (Bio Whittaker Inc.) and 10% (v/v) heat-inactivated fetal calf serum (Gibco). After reaching near-confluence, cells were cultured overnight in serum-free medium. Cells were subsequently stimulated for 15 min with indicated stimuli. Cellular lysates were prepared using lysis buffer (0.5% [v/v] Triton X-100, 0.1 M Tris–HCl pH 7.4, 100 mM NaCl, 1 mM MgCl2, 1 mM Na2VO4, mini complete protease inhibitor cocktail (Boeringer Mannheim, Roche, Basel, Switzerland)). Samples were subjected to SDS-PAGE on a 10% glycin-based gel. Resolved proteins were transferred to a polyvinylidene difluoride (PVDF) membrane, and non-specific binding sites were blocked by PBS/0.05% Tween-20/1% casein. The blots were incubated with rabbit polyclonal antibodies against phosphorylated ERK-1/2 (New England Biolabs, Beverly, MA), and secondary horseradish peroxidase conjugated anti-rabbit IgG antibodies. The enhanced chemoluminescent (ECL) Western blotting detection system (Amersham Pharmacia Biotech, Uppsala, Sweden) was used to reveal immunoreactivity.

2.6. Chemotaxis assay

Neutrophils were isolated from peripheral blood from healthy volunteers using Percoll density gradient centrifugation (density: 1.082 g/ml) as previously described [44]. Cells were resuspended at a concentration of 2.5 × 10^6 cells/ml in assay buffer (20 mM N-2-hydroxyethylpiperazine-N’-2-ethanesulfonic acid [Hepes buffer; pH 7.4], 132 mM NaCl, 6 mM KCl, 1.2 mM KH2PO4, 1 mM MgSO4, 5.5 mM glucose, 0.1 mM CaCl2 and 0.5% human serum albumin [Central Laboratory of the Netherlands Red Cross Blood Transfusion Service (CLB), Amsterdam, The Netherlands]) diluted 1:1 with serum-free RPMI supplemented with 2 mM l-glutamine, 20 U/ml penicillin, and 20 U/ml streptomycin.

Neutrophil chemotactic activity of peptides was assessed using a modified Boyden Chamber technique [19]. Briefly, 26 μl of chemotactic stimuli in assay buffer was added to the wells of the lower compartment and 50 μl of neutrophil suspension was added to the upper compartment of a 48-well chamber (Neuroprobe, Cabin John, MD, USA). Two filters separated the
compartments: a lower filter with a pore size of 0.45 μm (Millipore Products, Bedford, MA) and an upper filter with a pore size of 8 μm (Sartorius Filter, San Francisco, CA). After incubation for 90 min at 37 °C, the upper filters were removed, fixed in ethanol:butanol (80:20, v/v), and stained with Weigert solution. To determine neutrophil chemotactic activity, migrated neutrophils were counted in six random fields as a positive control chemotactic activity assay was conducted with different concentrations of the highest three levels of P60.4-Ac and LL-37 to compare their effects on the bacterial or yeast strain C. albicans was prepared as described above. The fungus growth. Antibacterial activity was examined using log-phase cultured bacteria in Trypticase Soy Broth at 37 °C. The cultures were diluted with 10 mM sodium phosphate buffer pH 7.4 to give approximately 5.0 × 10⁶ CFU/ml. Ten microliters of the diluted test strain was transferred to a 96-well plate and 100 μl of the different peptide concentrations diluted in 1% Trypticase Soy Broth in 10 mM Sodium Phosphate buffer pH 7.4 was added to each well. The plates were incubated at 37 °C for 24 h and are then scored for growth by visual inspection on a light box. They are then returned for incubation for a further 24 h after which time they are re-evaluated for growth. The yeast strain C. albicans was prepared as described above. The filamentous fungi A. niger was used as a spore suspension, cultured on Sabouraud's Dextrose Agar plates at 20–25 °C for 6–10 days or until adequate sporulation has occurred. The spores were harvested by scraping and if necessary the concentration was adjusted to a final concentration of 5 × 10⁶ CFU/ml. 2.7.2. Toxicity

We performed three other toxicity studies with P60.4-Ac in animals. In a primary skin irritation/corrosion study three rabbits were exposed to 0.5 ml phosphate buffered (20 mM, pH 7.5, isotonic with 0.9% NaCl solution) peptide solution (2 mg/ml), applied onto clipped skin for 4 h using a semi-occlusive dressing. Observations were made 1, 24, 48, and 72 h after exposure. Single samples of 0.1 ml of phosphate buffered (pH 7.5) peptide solution (2 mg/ml) were instilled into one eye of each of three rabbits to perform an acute eye irritation/corrosion study. Observations were made 1, 24, 48, and 72 h after instillation. Finally, the toxic potential of P60.4-Ac was assessed in a single and repeated dose toxicity study in rats. The study was performed based on the ICH Harmonised Tripartite Guideline: Non Clinical Safety Studies for the Conduct of Human Clinical Trials for Pharmaceuticals, 16 July 1997. For this reason, the peptide was administered daily intravenously in escalating doses. In this phase, the maximum tolerated dose (MTD) was established. Repeated dose toxicity was also studied in the MTD phase. In the dose escalation phase nine rats were divided in three groups and received 0.4, 2, or 8 mg/(kg day) for 2 days. Clinical signs were recorded twice daily on days of dosing and 1 day after dosing, body weights were recorded prior to the first dose and 1 day after dosing. In the MTD phase, five female and five male rats received 8 mg/(kg day) for 5 consecutive days. Clinical signs were recorded twice daily on days of dosing, body weight on days 1 and 6. Clinical laboratory investigations were performed prior to necropsy. Macroscopy was performed at termination of the MTD phase. 2.8. Antimicrobial assays

The in vitro antibacterial and antifungal activity of P60.4-Ac and LL-37 were determined as the minimum inhibitory concentrations (MIC) by a microdilution susceptibility test in 96-well microtiter plates, according to a modified version of Hancock's "Modified MIC method for cationic antimicrobial peptides" [13]. The antibacterial activity was tested on the reference strains Escherichia coli ATCC 8739, Pseudomonas aeruginosa ATCC 9027. The antifungal activity was evaluated on Candida albicans ATCC 10231 and Aspergillus niger ATCC 14406. The antimicrobial activity assay was conducted with different concentrations of P60.4Ac and LL-37 to compare their effects on the bacterial or fungal growth. Antibacterial activity was examined using log-phase cultured bacteria in Trypticase Soy Broth at 37 °C. The cultures were diluted with 10 mM sodium phosphate buffer pH 7.4 to give approximately 5.0 × 10⁶ CFU/ml. Ten microliters of the diluted test strain was transferred to a 96-well plate and 100 μl of the different peptide concentrations diluted in 1% Trypticase Soy Broth in 10 mM Sodium Phosphate buffer pH 7.4 was added to each well. The plates were incubated at 37 °C for 24 h and are then scored for growth by visual inspection on a light box. They are then returned for incubation for a further 24 h after which time they are re-evaluated for growth. The yeast strain C. albicans was prepared as described above. The filamentous fungi A. niger was used as a spore suspension, cultured on Sabourauds Dextrose Agar plates at 20–25 °C for 6–10 days or until adequate sporulation has occurred. The spores were harvested by scraping and if necessary the concentration was adjusted to a final concentration of 5 × 10⁶ CFU/ml. 2.9. Statistical analysis

LPS and LTA neutralization data are shown as mean ± standard deviation (S.D.). Statistical significance was determined by one-way ANOVA with the Statistical Package for Social Sciences
Q10 (neutral/polar) was replaced by E in an attempt to stabilize sequence RPLR was predicted to yield better felicity. In addition, Replacement of the C-terminal part of P60 (PRTE) by the might be an indication that the C-terminus of P60 is not helical.

The helix by possible salt bridges with neighboring positive

Prism.Differencesat

in LPS neutralization with a LAL assay and a whole blood (WB)

Peptides P60 and P60.4 were further examined for their efficacy in LPS neutralization with a LAL assay and a whole blood (WB) assay. Both assays showed comparable results. The peptide concentration whereby 50% of the LPS activity is inhibited is a measure of the peptide’s activity (IC50); a low IC50 indicates high potency of LPS neutralization. When the 50%-LPS inhibition results are compared, it is clear that LL-37 has the strongest activity with an IC50 of 0.29 ± 0.06 μM in the LAL assay and an IC50 of 0.27 ± 0.05 μM in the WB assay (Table 2). The activity of P60 did not differ significantly from LL-37. Peptide P60.4 showed significantly (P < 0.05) lower activity in the WB assay but was not significantly different from LL-37 in the LAL assay. In Fig. 2, the LPS inhibition curves determined with the LAL assay for LL-

3.1. Synthetic peptides

All different peptides were analyzed first in the LAL assay. The activity of all the different peptides is presented in Table 1 as 50% LPS neutralizing values in μM (IC50). It was obvious that the LPS neutralizing activity of LL-37 resides primarily in the C-terminal part of the peptide. Subsequently, we tried to further improve activity by amino acid modifications. Computer analysis (Peptide Companion) showed that P60 is likely to fold into an amphipatic helix, although the amphipatic structure is not ideal. Especially, P21 and T23 seem to be “misplaced”, which might be an indication that the C-terminus of P60 is not helical. Replacement of the C-terminal part of P60 (PRTE) by the sequence RPLR was predicted to yield better felicity. In addition, Q10 (neutral/polar) was replaced by E in an attempt to stabilize the helix by possible salt bridges with neighboring positive charges. D14 was replaced by R to increase the overall charge of the polar part of the helix. The effects of these modifications were investigated by evaluating the LPS neutralizing activity of the variants as indicated in Table 1. Based on these results, we selected the peptides P60 and P60.4 for further detailed investigation. An acetylated and amidated variant of P60 was made (P60-Ac) to improve stability and tested to investigate whether this would influence the properties of the peptide. Helical wheel presentations show the optimization of the predicted amphipatic α-helical structure of P60.4 resulting from amino acid substitutions in P60 (Fig. 1).

3.2. LPS and LTA neutralization

Peptides P60 and P60.4 were further examined for their efficacy in LPS neutralization with a LAL assay and a whole blood (WB) assay. To determine a significant response in the T cell proliferation assay, counts per minute in the presence of stimulus were compared to the relevant medium control using one-way ANOVA (SPSS and Graph Pad Prism). Elispot responses were analyzed with the Mann–Whitney U-test in SPSS and Graph Pad Prism. Differences at P-values <0.05 were considered significant.

To explore possible immune stimulatory or pro-inflammatory activity we performed several experiments with our selected

Table 2 – LTA and LPS neutralization ± S.D. for the different peptides

<table>
<thead>
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<th>Peptide</th>
<th>50%-LPS inhibition (μM)</th>
<th>50%-LTA inhibition (μM), n=3</th>
</tr>
</thead>
<tbody>
<tr>
<td>LL-37</td>
<td>0.29 ± 0.06 (n=5)</td>
<td>0.36 ± 0.10</td>
</tr>
<tr>
<td>P60</td>
<td>0.44 ± 0.14 (n=5)</td>
<td>0.62 ± 0.22</td>
</tr>
<tr>
<td>P60.4</td>
<td>0.51 ± 0.18 (n=5)</td>
<td>0.59 ± 0.38</td>
</tr>
<tr>
<td>P60-Ac</td>
<td>0.53 ± 0.24 (n=5)</td>
<td>0.70 ± 0.15 (n=2)</td>
</tr>
</tbody>
</table>

* Number of experiments.

b Significantly lower activity than LL-37 (P < 0.05).

c Significantly lower activity than P60 (P < 0.05).

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</tr>
</tbody>
</table>

To investigate whether acetylation and amidation would influence the activity of a synthetic peptide, P60-Ac was also tested for LPS and LTA neutralization. Except from the WB assay for LPS neutralization P60-Ac did not differ significantly from P60. It appeared that stabilizing a peptide by acetylation and amidation did not influence the LPS and LTA neutralizing capacity to a large extend (Table 2; Fig. 2).

3.3. Immunogenicity

To explore possible immune stimulatory or pro-inflammatory activity we performed several experiments with our selected...
3.3.1. T cell assays

The two selected synthetic peptides P60 and P60.4 were tested in an Elispot to assess cytokine production and in a T cell proliferation assay. For analysis of the T cell proliferation IL-2 served as a positive control and increased the mean T cell proliferation significantly \( (P < 0.01) \) (Fig. 3). A concentration of 1 or 10 \( \mu \text{g/ml} \) peptide did not induce any significant stimulation of the T cells. At 100 \( \mu \text{g/ml} \) LL-37 induced a significantly increased mean T cell proliferation \( (P < 0.05) \) whereas, P60 and P60.4 did not induce significant T cell proliferation. The synthetic peptide P60.4 resulted in a significantly \( (P < 0.05) \) lower T cell proliferation compared to LL-37. When donors were compared individually, LL-37 induced significant \( (P < 0.05) \) increased T cell proliferation in six out of eight donors, P60 in seven out of eight donors, and P60.4 in only two out of eight donors (Table 3). The T cell proliferation of P60-Ac was not significantly different from the medium (Fig. 3).

The Elispot assay was used to evaluate the effect of our peptides on cytokine production by PBMC, thereby providing data on their putative modulating effects on immune function in vivo [43]. Compared with medium alone, no significant IFN-\( \gamma \) responses were detected for the peptides (Fig. 4A). When the donors were analyzed individually, LL-37 induced significant \( (P < 0.05) \) increase in IFN-\( \gamma \) producing T cells in four out of eight donors (Table 3). P60 induced a significant increase in one out of eight donors and P60.4 in two out of eight donors. No significant IL-10 responses were detected for the peptides when compared with medium alone (Fig. 4B). LL-37 and P60 both induced a significant \( (P < 0.05) \) increase in IL-10 producing cells in four out of eight donors (Table 3); whereas, P60.4 induced a significant \( (P < 0.05) \) response in only two out of eight donors. Stimulation with P60-Ac did not induce significantly different mean values for IFN-\( \gamma \) producing T cells and IL-10 producing cells compared to the other peptides or compared to medium alone.

3.3.2. Chemotaxis assay

Cellular migration of neutrophils due to chemotaxis induced by our selected synthetic peptides is represented in Fig. 5 and Table 3. The positive control, FMLP, induced the highest cellular migration.

**Table 3 - Summary of the immunogenicity experiments with the synthetic peptides compared to LL-37**

<table>
<thead>
<tr>
<th>Compound</th>
<th>( \gamma )-IFN Elispot(^a)</th>
<th>IL-10 Elispot(^a)</th>
<th>T cell proliferation(^a)</th>
<th>ERK-activation(^b)</th>
<th>Chemotaxis(^c)</th>
</tr>
</thead>
<tbody>
<tr>
<td>P60</td>
<td>1/8</td>
<td>4/8</td>
<td>7/8</td>
<td>–</td>
<td>76 ± 39 (n = 3)</td>
</tr>
<tr>
<td>P60.4</td>
<td>3/8</td>
<td>2/8</td>
<td>2/8</td>
<td>±</td>
<td>0 ± 0 (n = 2)</td>
</tr>
<tr>
<td>LL-37</td>
<td>4/8</td>
<td>4/8</td>
<td>6/8</td>
<td>+</td>
<td>84 ± 17 (n = 5)</td>
</tr>
<tr>
<td>P60-Ac</td>
<td>3/8</td>
<td>4/8</td>
<td>4/8</td>
<td>±</td>
<td>61 ± 36 (n = 4)</td>
</tr>
</tbody>
</table>

\(^a\) Results are shown as number of individual donors that induced a significant \( (P < 0.05) \) increase when peptide-stimulated responses were compared to medium induced responses.

\(^b\) At 7.3 \( \mu \text{M} \), experiment performed in duplicate.

\(^c\) Values represent migration of neutrophil cells counted per six high-power fields.

\(^d\) Significantly \( (P < 0.05) \) lower than LL-37 induced migration.
migration. LL-37 and P60 induced a significant (P < 0.05) migration of neutrophils compared to medium induced migration, whereas P60.4 did not induce any migration. Acetylation and amidation of P60 induced no significant change in chemotaxis.

3.3.3. ERK1/2-activation
Possible activation of epithelial cells by the peptides was examined in the extracellular signal-regulated kinase (ERK)-activation assay. ERK1/2 is part of the MAP-kinase signaling pathway, which has been shown to be involved in various cellular processes, including proliferation, differentiation, cell survival and expression of genes encoding pro-inflammatory mediators like cytokines [1]. None of the peptides, including LL-37, induced ERK-activation at a concentration of 2.4 μM (Fig. 6). At 7.3 μM, LL-37 induced marked phosphorylation of ERK1/2, indicative of epithelial cell activation. P60.4 induced a moderate ERK-activation at this concentration, whereas P60 did not induce clear ERK-activation. From the original Western blot and the accompanying intensity score of the ERK-activation (Table 3) it is clear that at a concentration of 7.3 μM our synthetic peptides P60 and P60.4 induced a lower ERK-activation than LL-37. P60-Ac induced a moderate ERK-activation, which was higher than the P60 induced ERK-activation.

3.4. In vivo toxicity studies
P60.4 showed the most favorable results in case of immunogenicity and LL-37 comparable LPS and LTA neutralization. Since P60 and P60-Ac did not differ significantly in activity or in immunogenicity, this is indicating that acetylation and amidation of the peptides does not affect their activity in vitro. Because acetylation and amidation increases stability, and because P60.4 caused less T cell and neutrophil activation, toxicity studies were performed with P60.4-Ac.

3.4.1. Ototoxicity
Ototoxicity was measured in guinea pigs. Round window application of PBS, which was used as a control, did not result in a threshold change at 22 days after surgery. Formulation buffer resulted in a threshold change of 2 dB after 22 days. Cisplatin, on the other hand, induced threshold changes of, respectively, −49 and −64 dB, which indicate a severe hearing loss (Table 4). This part of the experiments served as a positive control for the ototoxicity study. Peptide P60.4-Ac (2 mg/ml) in PBS induced a threshold change of −7 dB 22 days after surgery. Both animals that received P60.4-Ac in formulation buffer produced a threshold change of 1 dB.

### Table 4 – Ototoxicity measured in guinea pigs

<table>
<thead>
<tr>
<th>Group 1</th>
<th>PBS</th>
<th>Cisplatin</th>
<th>P60.4-Ac&lt;sup&gt;a&lt;/sup&gt;</th>
<th>Formulation&lt;sup&gt;b&lt;/sup&gt;</th>
<th>P60.4-Ac&lt;sup&gt;c&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>Post-surgery</td>
<td>1</td>
<td>−2</td>
<td>−1</td>
<td>−2</td>
<td>1</td>
</tr>
<tr>
<td>3 Days</td>
<td>−3</td>
<td>−32</td>
<td>−38</td>
<td>−2</td>
<td>−8</td>
</tr>
<tr>
<td>7 Days</td>
<td>−3</td>
<td>−32</td>
<td>−45</td>
<td>0</td>
<td>−33&lt;sup&gt;d&lt;/sup&gt;</td>
</tr>
<tr>
<td>14 Days</td>
<td>0</td>
<td>−30</td>
<td>−59</td>
<td>12</td>
<td>−1</td>
</tr>
<tr>
<td>22 Days</td>
<td>0</td>
<td>−49</td>
<td>−64</td>
<td>−7</td>
<td>2</td>
</tr>
</tbody>
</table>

Values represent Δpresurgery in dB.

<sup>a</sup> Solution in PBS.

<sup>b</sup> 7% Macrogol10.000 in isotone [NaCl] preserved [0.02% benzalkonium chloride and 0.1% Na₂EDTA] 20 mM phosphate pH 5.5.

<sup>c</sup> Solution in formulation buffer.

<sup>d</sup> Unreliable measurements due to bad wire.
The antimicrobial activity of P60.4-Ac was evaluated against two Gram-negative strains and against the fungi C. albicans and A. niger and compared with LL-37. The MIC values for each peptide are reported in Table 5. P60.4-Ac showed a higher or equal activity against the Gram-negative strains E. coli and P. aeruginosa compared to LL-37. In some cases, bactericidal activity was also determined for both peptides. P60.4-Ac showed an MIC at 6 μM against C. albicans and may well be fungicidal at 18 μM. P60.4-Ac at 18 μM inhibited the germination of A. niger spores for 24 h, whereas LL-37 shows no activity against A. niger.

### 3.4.2. General toxicity

No skin irritation was caused by 4 h exposure to phosphate buffered peptide solution of P60.4-Ac (data not shown). Instillation of the peptide solution resulted in redness of the conjunctivae that resolved completely within 24 h after instillation (data not shown). No mortality occurred in the dose escalation study with 0.4, 2, and 8 mg/(kg day). Furthermore, no clear deviations were noted in clinical signs and body weight. During the maximum tolerated dosage (MTD) phase also no mortality occurred and no clear peptide related findings were noted in clinical signs, body weight, hematology and clinical biochemistry parameters and at macroscopic examination. An intravenous MTD of peptide P60.4-Ac could not be determined but will be higher than 8 mg/kg.

### 3.5. Antimicrobial activity

The antimicrobial activity of P60.4-Ac was evaluated against two Gram-negative strains and against the fungi C. albicans and A. niger and compared with LL-37. The MIC values for each peptide are reported in Table 5. P60.4-Ac showed a higher or equal activity against the Gram-negative strains E. coli and P. aeruginosa compared to LL-37. In some cases, bactericidal activity was also determined for both peptides. P60.4-Ac showed an MIC at 6 μM against C. albicans and may well be fungicidal at 18 μM. P60.4-Ac at 18 μM inhibited the germination of A. niger spores for 24 h, whereas LL-37 shows no activity against A. niger.

### 4. Discussion

In this study, we have developed synthetic derivatives of the human antimicrobial peptide hCAP18/LL-37. By downsizing the LL-37 peptide to 22-mer peptides we identified four peptides with good LPS neutralizing activity. To increase the activity, six elongated variants of these four peptides were made of which peptide variant P60 displayed the most powerful LPS neutralizing activity. Subsequently, modifications of this peptide were made in order to increase the hydrophobicity and cationicity, which resulted in peptide P60.4. The LPS and LTA neutralizing capacities of P60 and P60.4 were further analyzed in a LAL assay and a whole blood assay and compared with LL-37. We found a similar neutralization capacity of the two selected peptides when compared to endogenous LL-37.

Several studies showed that the C-terminal domain of hCAP18 could neutralize various activities of LPS [20,24,34] and LTA [34]. Nagaoka et al. [24] identified an 18-mer peptide derived from hCAP18/LL-37, which is also included in our 22-mer peptide. By modifying hydrophobicity and cationicity of this peptide they could augment LPS neutralizing activities of this peptide. In contrast to the cytotoxic activity of the 18-mer peptide described by Nagaoka et al., our peptide appeared to show less cytotoxic activity. It needs to be noted, however, that the peptides from the study of Nagaoka et al. and our peptides were evaluated in different assays.

When biologically active peptides are used clinically in their natural form, their biologic effects are often rapidly lost in vivo due to rapid proteolytic degradation of the active form of the peptide [21]. There is evidence that this cleavage of short peptides may be markedly inhibited by modifications of the N- and C-terminal residues [30]. Modifications like acetylation or amidation on the other hand, can also inhibit or reduce the activity of peptides [29,36]. To improve stability, we synthesized P60-Ac, the N-terminally acetylated and C-terminally amidated variant of P60. Our results showed that the LPS and LTA neutralization activity of the acetylated and amidated peptide as well as the immunogenicity of this peptide was not markedly changed.

Cytokines are direct mediators of inflammation and influence the progress and direction of many immunological reactions. Perturbation of the balance in cytokine production is widely recognized as a critical factor in several disease states. This balance is already disturbed in the case of OME [37] and sinusitis [46]. Various antimicrobial peptides have been shown to display activities related to inflammation and induction of immunity, which are distinct from their antimicrobial and LPS neutralizing activities [18]. LL-37 is known to stimulate epithelial cells [41] and macrophages [34]. Moreover, LL-37 is chemotactic for T cells [9] but until now it has not yet been demonstrated that LL-37 can also stimulate T cell proliferation and cytokine production. In the present study, we demonstrate that LL-37 increases T cell proliferation and cytokine production. Furthermore, we demonstrated that P60 and P60.4 induce cytokine production by T cells to an equal or lower level than the endogenous LL-37 peptide. Based on the nature of the Elispot assay, we cannot rule out the possibility that IL-10 was derived from monocytes. In the T cell proliferation assay, we observed the lowest response after incubation with our peptide P60.4. The
results indicate a lower stimulation of T cells by our synthetic developed peptides when compared to LL-37. We assume that this low T cell reactivity may be favorable because this might limit T cell activation in vivo.

For in vivo use of the selected LPS neutralizing peptides in upper airway infections, it is also important to explore the effects of the peptides on activation of epithelial cells. Therefore, we studied the effects of the peptides on phosphorylation of extracellular signal-regulated kinase (ERK1/2), in NCI-H292 bronchial epithelial cells. Tjabringa et al. [41] demonstrated that LL-37 activates airway epithelial cells by activating ERK1/2. Our selected synthetic peptides induced a lower activation of ERK1/2 in epithelial cells when compared to endogenous LL-37. Since excessive epithelial stimulation may lead to changes in epithelial phenotype (squamous and mucus differentiation) and inflammation, we assume that an ideal peptide should have maximal LPS neutralizing capacity and minimal stimulatory effects on epithelial cells. Therefore, it appears that our selected synthetic peptides, P60 and P60.4, may be suitable for therapeutic application in upper airway infections.

LL-37 is known to be chemotactic to neutrophils, monocytes, T cells, and mast cells [9,27]. We demonstrated a comparable chemotactic activity towards neutrophils for P60 and for LL-37. Remarkably, in this assay P60.4 did not show any chemotactic activity. These data suggest that the amino acid substitutions in P60.4 resulted in a decreased interaction with the formyl peptide receptor employed by LL-37 to exert its chemotactic activity [9]. Recruitment of inflammatory cells to sites of microbial invasion may be beneficial during acute infection. However, for treatment of chronic infections it is more favorable to dampen the immune response and not to attract more inflammatory cells. This consideration makes peptide P60.4 a very attractive candidate for further development.

Since the developed peptide was designed for use in treatment of upper respiratory tract infections, we also tested the possible otoxicity of P60.4-Ac in a formulation solution. Some studies have demonstrated that common topical otomicrobial agents can induce cochlear hair cell damage [4,32]. It is also known that neutrophil cationic peptides and cationic antibiotics such as polymyxin B have a limited clinical use because of their potential toxicity [33,38]. We could demonstrate that P60.4-Ac does not induce any signs of possible otoxicity. Additionally, we found that repeated intravenous administration of peptide P60.4-Ac to male and female rats at 8 mg/(kg day) on 5 consecutive days did not result in clear peptide related changes in clinical signs, body weight, hematology, clinical biochemistry, and macroscopic findings. Furthermore, no skin or lasting eye irritation was observed in the acute skin and eye irritation/corrosion study. So far, the results of these toxicity experiments indicate that our peptide can be used safely in further studies.

We tested the antimicrobial activity of P60.4-Ac against E. coli, P. aeruginosa, C. albicans, and A. niger. Our designed peptide P60.4-Ac showed a higher or equal antimicrobial activity than LL-37 against all the strains tested. When compared to LL-37 it seems that our designed antimicrobial peptide shows a shift in biological activity toward antimicrobial function and away from the ability to stimulate a host response. Murakami et al. [23] have also revealed this shift. They showed that human sweat contains shorter variants of LL-37 generated by proteolytic activity in sweat. Some of these peptides showed similar or enhanced antimicrobial action when compared to LL-37, but a decreased ability to stimulate IL-8 release from keratinocytes. In addition, predicted processing variants of the rat homologue of LL-37, rCRAMP, showed a similar swift in activity [40].

For treatment of upper airway infections antibiotics are still most frequently used, but over the past decade levels of bacterial resistance to antibiotics have risen dramatically [22]. As a result, there is a growing need to discover and introduce new drugs. Besides their direct antimicrobial function, antimicrobial peptides play an important role in the protection of epithelial surfaces through other activities. They have multiple roles as mediators of inflammation with impact on epithelial and inflammatory cells influencing diverse processes such as cell proliferation, immune induction, wound healing, cytokine release, and chemotaxis [18]. Antimicrobial peptides are attractive candidates for clinical development because of their selectivity, their speed of action and because bacteria may not easily develop resistance against them [5]. Furthermore, unlike many defensins, several of the cathelicidin-derived peptides retain broad-spectrum activity at physiologic or elevated salt concentrations [42]. However, the use of antimicrobial peptides can be limited by their non-selective toxicity, low stability, immunogenicity, and their costs of production.

In this study, we have developed synthetic derivatives of LL-37. One peptide, P60.4, which showed similar LPS and LTA neutralizing activity but with lower T cell stimulation, lower epithelial cell activation, and with much lower chemotactic capacity when compared to LL-37, was selected. In addition, a more stable acetylated and amidated form this peptide showed no signs of toxicity and had an improved antimicrobial activity when compared to LL-37. Therefore, our synthetic peptide may have therapeutic potential for bacterial and fungal infections and has potential as new therapeutic agent for upper respiratory tract infections.

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