High Cerebrospinal Fluid (CSF) Penetration and Potent Bactericidal Activity in CSF of NZ2114, a Novel Plectasin Variant, during Experimental Pneumococcal Meningitis

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Plectasin is the first defensin-type antimicrobial peptide isolated from a fungus and has potent activity against gram-positive bacteria. By using an experimental meningitis model, the penetration of plectasin into the cerebrospinal fluid (CSF) of infected and uninfected rabbits and the bactericidal activities in CSF of the plectasin variant NZ2114 and ceftriaxone against a penicillin-resistant Streptococcus pneumoniae strain (NZ2114 and ceftriaxone MICs, 0.25 and 0.5 μg/ml, respectively) were studied. Pharmacokinetic analysis showed that there was a significantly higher level of CSF penetration of NZ2114 through inflamed than through noninflamed meninges (area under the concentration-time curve for CSF/area under the concentration-time curve for serum, 33% and 1.1%, respectively; P = 0.03). The peak concentrations of NZ2114 in purulent CSF were observed --3 h after the infusion of an intravenous bolus of either 20 or 40 mg/kg of body weight and exceeded the MIC >10-fold for a 6-h study period. Treatment with NZ2114 (40 and 20 mg/kg at 0 and 5 h, respectively; n = 11) caused a significantly higher reduction in CSF bacterial concentrations than therapy with ceftriaxone (125 mg/kg at 0 h; n = 7) at 3 h (median changes, 3.7 log_{10} CFU/ml [interquartile range, 2.5 to 4.6 log_{10} CFU/ml] and 2.1 log_{10} CFU/ml [interquartile range, 1.7 to 2.6 log_{10} CFU/ml], respectively; P = 0.001), 5 h (median changes, 5.2 log_{10} CFU/ml [interquartile range, 3.6 to 6.1 log_{10} CFU/ml] and 3.1 log_{10} CFU/ml [interquartile range, 2.6 to 3.7 log_{10} CFU/ml], respectively; P = 0.01), and 10 h (median changes, 5.6 log_{10} CFU/ml [interquartile range, 5.2 to 5.9 log_{10} CFU/ml] and 4.2 log_{10} CFU/ml [interquartile range, 3.6 to 5.0 log_{10} CFU/ml], respectively; P = 0.03) after the start of therapy as well compared to the CSF bacterial concentrations in untreated rabbits with meningitis (n = 7, P < 0.05). Also, significantly more rabbits had sterile CSF at 5 and 10 h when they were treated with NZ2114 than when they were treated with ceftriaxone (67% [six of nine rabbits] and 0% [zero of seven rabbits], respectively, at 5 h and 75% [six of eight rabbits] and 14% [one of seven rabbits], respectively, at 10 h; P < 0.05). Due to its excellent CSF penetration and potent bactericidal activity in CSF, the plectasin variant NZ2114 could be a promising new option for the treatment of CNS infections caused by gram-positive bacteria, including penicillin-resistant pneumococcal meningitis.

Streptococcus pneumoniae is the leading cause of bacterial meningitis, and pneumococcal meningitis still remains a disease with high rates of mortality and morbidity (8). The cornerstone of treatment for bacterial meningitis is the prompt initiation of adequate antibiotic therapy (12), and the rapid sterilization of the cerebrospinal fluid (CSF) with antibiotic initiation of adequate antibiotic therapy (12), and the rapid sterilization of the cerebrospinal fluid (CSF) with antibiotic therapy has been associated with a better clinical outcome (4). The worldwide emergence of antibiotic-resistant strains of S. pneumoniae still demands the development of new antimicrobial agents for the treatment of pneumococcal disease, including meningitis.

Antimicrobial peptides are evolutionarily ancient molecules with antimicrobial effects that are widely distributed in plants and animals and that are used by the innate immune system to control infections. They are a new class of antimicrobial agents, but their mechanisms of action have not yet been determined. Most antimicrobial peptides are cationic and readily bind to the negatively charged bacterial membrane. Their activity may be promoted by their ability to compromise bacterial membranes (e.g., by fatal depolarization of the bacterial membrane, degradation of the cell wall by hydrolases, or disturbance of membrane function), or they may interact with critical intracellular targets (for a review, see reference 13).

Plectasin was the first defensin-like antimicrobial peptide isolated from a black saprophytic ascomycete (Pseudoplectania nigella) and showed potent activity against gram-positive bacteria in vitro and in vivo (5). NZ2114 was identified in a high-throughput mutation and screening campaign aimed at identifying variants of plectasin with improved potency against staphylococci and streptococci (10). NZ2114 exhibited improved in vitro activity compared with the activities of other plectasin variants against staphylococci, including Staphylococcus aureus, as well as S. pneumoniae and hemolytic streptococcal strains resistant to clinically used antibiotics (11). Little is known about the distribution of antimicrobial peptides into various body tissues, including their penetration across the blood-brain barrier, which forms a tight membrane that limits the entry of many antimicrobial agents into the central nervous system. Therefore, when a new antimicrobial agent is introduced, it is important to study its penetration into the brain.
and to determine the bactericidal activity of the drug in this compartment. Correspondingly, we studied the CSF penetration of NZ2114 as well as its bactericidal activity in CSF against a penicillin-resistant *S. pneumoniae* strain in an experimental meningitis model.

**MATERIALS AND METHODS**

**Test organism.** A penicillin-resistant but ceftriaxone-sensitive *Streptococcus pneumoniae* type 9V strain originally isolated from the CSF of a 78-year-old woman was used for all meningitis experiments. The MICs of NZ2114, ceftriaxone, and penicillin were 0.25 mg/liter, 0.5 mg/liter, and 1 mg/liter, respectively, as determined by the microdilution broth method according to the guidelines of the CLSI (formerly the NCCLS) (6).

**Antimicrobial agents.** The NZ2114 plectasin variant was produced, purified, and formulated at Novozymes A/S, Bagsvaerd, Denmark. In short, *Aspergillus oryzae* MT3343 was selected as the production host, and the fermentation conditions were optimized at Novozymes. The fermented broth was pretreated, filtered, and centrifuged before the recovered product was further processed by two chromatographic purifications by use of a cation exchange and hydrophobic interaction, followed by filtration with a 30-kDa cutoff and final diafiltration. The active pharmaceutical ingredient was diluted into the final formulation buffer (10 mM sodium acetate, 0.9% NaCl, pH 5.0) after diafiltration and germ filtered before filled in vials.

Stock solutions were made by dissolving NZ2114 (batch PSE6012; purity, 95%) in potassium-natrium-glucose infusion liquid (06105C03; Fresenius Kabi, Uppsala, Sweden) to a final concentration of 18.3 g/liter (vehicle pH, 5). Ceftriaxone (no. C-5793; Sigma Chemical Co., St. Louis, MO) was dissolved in sterile water to a final concentration of 50 g/liter. The antibiotics were administered intravenously as a bolus infusion over 5 to 10 min.

**Determination of NZ2114 concentrations in CSF and serum.** The NZ2114 concentrations in CSF and serum were determined by high-pressure liquid chromatography. The lower limit of detection was 50 mg/liter.

**Determination of bacterial concentrations.** Bacterial concentrations in CSF were determined by plating 140 μl of undiluted CSF and 10-fold serial dilutions of CSF (in duplicate) on 5% horse blood agar plates (Statens Serum Institute, Copenhagen, Denmark). Thus, the lowest detectable bacterial count was 7 CFU/ml. The bacterial counts in different dilutions of CSF were compared to exclude the possibility of the presence of significant carryover phenomena.

**Meningitis model.** The experimental protocols were approved by the Danish Animal Experiments Inspectorate (Dyreforsøgstilsynet). A rabbit meningitis model previously described in detail was used (9). New Zealand White rabbits (weight, 2.5 to 3.0 kg) were anesthetized with midazolam (Dormicum; F. Hoffmann-La Roche AG, Basel, Switzerland) at 0.5 mg/kg of body weight subcutaneously and a combination of fentanyl-fluanisone (Hypnorm; Janssen Pharmaceutical, Neuss, Germany). Thus, the lowest detectable bacterial count was 7 CFU/ml. The bacterial counts in different dilutions of CSF were compared to exclude the possibility of the presence of significant carryover phenomena.

**Statistical analysis.** All results are provided as medians and interquartile ranges. For calculation of the reduction in CSF bacterial concentrations (e.g., the change in the log₁₀ CFU/ml from 0 to 3 h), a CSF concentration under the detection limit was assigned a value of 1 CFU/ml. Comparison between groups was performed by the nonparametric Mann-Whitney test for continuous data and by the Fisher exact test for categorical data. *P* values less than 0.05 were considered statistically significant.

**RESULTS**

Pharmacokinetics of NZ2114. CSF and serum concentration-time curves and the values of the pharmacokinetic indices after the administration of intravenous bolus infusions of NZ2114 are shown in Fig. 1 and Table 1, respectively. Whereas the distribution of NZ2114 in the systemic compartment was comparable for both infected and uninfected rabbits (one-compartment model with a *t₁/₂* of 54 min [interquartile range, 49 to 56 min]), the CSF penetration (ratio of the AUC for CSF/AUC for serum) was significantly higher in rabbits with meningitis than in the uninfected controls by use of the current dosing regimens (one dose of 20 and 40 mg/kg; Fig. 1A and B): 33% (interquartile range, 24 to 50%) and 1.1% (interquartile range, 0.7 to 1.9%), respectively (*P* = 0.03). The *Cₘₐₓ* was 20 to 30 times higher in rabbits with meningitis than in uninfected rabbits (e.g., 10.4 mg/liter [interquartile range, 8.8 to 11.7 mg/liter] and 0.3 mg/liter [interquartile range, 0.2 to 0.4 mg/liter], respectively, after administration of a 40-mg/kg bolus infusion) and was observed >3 h after the bolus infusion. The CSF concentration did not decrease significantly after the time to *Cₘₐₓ* (1/2 of 54 min) was considerably more than 10 times longer in the CSF than in serum) and remained above the MIC for 6 h study period for rabbits with meningitis, whereas it was below the MIC for the uninfected controls most of the time. During the treatment efficacy studies (40 and 20 mg/kg at 0 and 5 h, respectively), NZ2114 concentrations in CSF and blood remained at concentrations above 10× the MIC during the 10-h study period (Fig. 1C).

**Treatment efficacy of NZ2114 in rabbits with meningitis caused by a penicillin-resistant pneumococcal strain.** CSF bacterial concentrations are shown in Fig. 2. Before the initiation of antibiotic therapy (at 0 h), no significant differences in the CSF bacterial concentrations were observed among the three experimental groups: for NZ2114-treated rabbits (*n* = 7), 6.0 log₁₀ CFU/ml (interquartile range, 5.2 to 7.0 log₁₀ CFU/ml); for ceftriaxone-treated rabbits (*n* = 7), 6.2 log₁₀ CFU/ml (interquartile range, 5.3 to 6.7 log₁₀ CFU/ml); and for untreated rabbits with meningitis (*n* = 7), 6.7 log₁₀ CFU/ml (interquartile range, 6.2 to 7.1 log₁₀ CFU/ml) (*P* > 0.05). The NZ2114-treated rabbits had significantly lower CSF bacterial concentrations than the ceftriaxone-treated rabbits at 3 h (2.7
log_{10} CFU/ml [interquartile range, 0.8 to 3.1 log_{10} CFU/ml] and 4.2 [interquartile range, 3.1 to 5.0 log_{10} CFU/ml], respectively; \( P = 0.01 \), 5 h (0.8 log_{10} CFU/ml [interquartile range, 0.8 to 2.2 log_{10} CFU/ml] and 2.8 log_{10} CFU/ml [interquartile range, 2.6 to 3.2 log_{10} CFU/ml], respectively; \( P = 0.005 \)), and 10 h (0.8 log_{10} CFU/ml [0.8 to 1.1 log_{10} CFU/ml] and 1.3 log_{10} CFU/ml [1.0 to 2.9], respectively; \( P = 0.05 \)) after the start of antibiotic therapy, whereas antibiotic-treated rabbits had significantly lower CSF bacterial counts than untreated rabbits with meningitis at all three time points (\( P < 0.02 \)) (Fig. 2). This

FIG. 1. CSF and serum concentrations of NZ2114 in infected and uninfected rabbits. (A and B) NZ2114 was administered as an intravenous bolus infusion over 5 to 10 min at 0 h (arrows) (n = 2 for each group); (C) NZ2114 was administered at 0 h (40 mg/kg) and at 5 h (20 mg/kg) (arrows) (n = 11). Data are shown as medians and ranges (A and B) or interquartile ranges (C).
was a significantly higher reduction in the numbers of CFU for rabbits treated with NZ2114 than for rabbits treated with ceftriaxone: at 3 h, the reductions were 3.7 log<sub>10</sub> CFU/ml/h (interquartile range, 2.5 to 4.6 log<sub>10</sub> CFU/ml/h) and 2.1 log<sub>10</sub> CFU/ml/h (interquartile range, 1.7 to 2.6 log<sub>10</sub> CFU/ml/h), respectively (P<0.001); at 5 h, the reductions were 5.2 log<sub>10</sub> CFU/ml/h (interquartile range, 3.6 to 6.1 log<sub>10</sub> CFU/ml/h) and 3.1 log<sub>10</sub> CFU/ml/h (interquartile range, 2.6 to 3.7 log<sub>10</sub> CFU/ml/h), respectively (P = 0.01); and at 10 h, the reductions were 5.6 log<sub>10</sub> CFU/ml/h (interquartile range, 5.2 to 5.9 log<sub>10</sub> CFU/ml/h) and 4.2 log<sub>10</sub> CFU/ml/h (interquartile range, 3.6 to 5.0 log<sub>10</sub> CFU/ml/h), respectively (P = 0.03). In addition, significantly more NZ2114-treated rabbits than ceftriaxone-treated rabbits had sterile CSF: at 5 h, 67% (six of nine rabbits) and 0% (zero of seven rabbits), respectively, and at 10 h, 75% (six of eight rabbits) and 14% (one of seven rabbits), respectively (P < 0.05).

DISCUSSION

To our knowledge, this is the first study to describe the CSF penetration and bactericidal activity in CSF of an antimicrobial peptide. The plectasin variant NZ2114 had a CSF penetration of 33% through inflamed meninges, which resulted in CSF concentrations above the MICs for most gram-positive pathogens, including S. pneumoniae. Interestingly, this level of CSF penetration was higher than the levels for most other antimicrobial agents except fluoroquinolones (e.g., ceftriaxone, ~15%; vancomycin, 13%; moxifloxacin, 81%) (9). This indicates that even relatively large and water-soluble molecules like NZ2114 (a peptide of 40 amino acids in length) (5), which has a molecular mass (4.4 kDa) 10 times larger than the molecular masses of most other antimicrobial agents (e.g., the ceftriaxone molecular mass is 0.6 kDa) (7), readily enter the CSF when the blood-brain barrier is inflamed. In contrast, the level of CSF penetration of NZ2114 through an intact blood-brain barrier was poor (1.1%), and the concentration in CSF hardly reached the MIC after the administration of an intravenous dose of 40 mg/kg, but its level of penetration through noninflamed meninges was still comparable to that of ceftriaxone (2%) (1). Besides alterations in blood-brain barrier permeability, the lipid solubility, and the molecular mass of the drug, the level of protein binding and whether it is a substrate for P-glycoprotein should be considered.
of various active influx and efflux pumps of the brain endothelium also influence the CSF penetration of antibiotics (for a review, see reference 2) and peptides (for a review, see reference 3). The passage of NZ2114 across an intact blood-brain barrier in part showed saturable features, since the CSF penetration decreased with higher doses; however, this was not the case for an inflamed blood-brain barrier.

The bactericidal activity of NZ2114 against penicillin-resistant pneumococci in CSF was high, resulting in a rapid killing rate in CSF (~1 log_{10} CFU/ml/h) during experimental pneumococcal meningitis in rabbits, and this killing rate was significantly more rapid than that achieved with therapy with ceftriaxone, a recommended treatment for pneumococcal meningitis (12). Moreover, sterilization of the CSF was rapid and was observed for the majority of rabbits within 5 h of therapy with NZ2114, and no regrowth was observed during the following 5 h of therapy. A previous study investigated the in vivo activity of the parent wild-type molecule against pneumococci by using a mouse peritonitis model and a mouse pneumonia model and found, in accordance with the findings of the present study, that plectasin had excellent killing rates that were comparable to those achieved with therapy with other antibiotics (5).

Most antimicrobial peptides are positively charged and readily bind to the negatively charged phospholipid of the bacterial cytoplasmic membrane, which promotes the rapid killing of microorganisms at concentrations equivalent or close to the MIC. In addition, contrary to the case when conventional antibiotics are used, the development of resistance to antimicrobial peptides seems to be surprisingly rare. It has been hypothesized that antimicrobial peptides primarily work by compromising bacterial membranes (e.g., by fatal depolarization of the bacterial membrane, degradation of the cell wall by hydrolyse, and the disturbance of membrane functions [for a review, see reference 13]). However, preliminary results showed that the mechanism by which NZ2114 kills bacteria is different from membrane lysis (H.-H. Kristensen, personal communication). In addition, no cross-resistance to other classes of antibiotics has yet been observed (Kristensen, personal communication). Thus, due to the emergence of resistant pneumococci and the potent bactericidal activity of NZ2114 in CSF, NZ2114 could be an interesting option for the treatment of pneumococcal meningitis and warrants further clinical and experimental evaluations.

In conclusion, NZ2114 could be a promising new option for the treatment of central nervous system infections, including penicillin-resistant pneumococcal meningitis, due to its excellent CSF penetration and potent bactericidal activity in CSF.

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REFERENCES