

Disulfide cross-linking influences symbiotic activities of nodule peptide NCR247

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Interactions of rhizobia with legumes establish the chronic intracellular infection that underlies symbiosis. Within nodules of inverted repeat-lacking clade (IRLC) legumes, rhizobia differentiate into nitrogen-fixing bacteroids. This terminal differentiation is driven by host nodule-specific cysteine-rich (NCR) peptides that orchestrate the adaptation of free-living bacteria into intracellular residents. *Medicago truncatula* encodes a family of >700 NCR peptides that have conserved cysteine motifs. NCR247 is a cationic peptide with four cysteines that can form two intramolecular disulfide bonds in the oxidized forms. This peptide affects *Sinorhizobium meliloti* transcription, translation, and cell division at low concentrations and is antimicrobial at higher concentrations. By preparing the three possible disulfide-cross-linked NCR247 regioisomers, the reduced peptide, and a variant lacking cysteines, we performed a systematic study of the effects of intramolecular disulfide cross-linking and cysteines on the activities of an NCR peptide. The relative activities of the five NCR247 variants differed strikingly among the various bioassays, suggesting that the NCR peptide-based language used by plants to control the development of their bacterial partners during symbiosis is even greater than previously recognized. These patterns indicate that certain NCR bioactivities require cysteines whereas others do not. The results also suggest that NCR247 may exert some of its effects within the cell envelope whereas other activities occur in the cytoplasm. BacA, a membrane protein that is critical for symbiosis, provides protection against all bactericidal forms of NCR247. Oxidative folding protects NCR247 from degradation by the symbiotically relevant metalloprotease HrrP (host range restriction peptidase), suggesting that disulfide bond formation may additionally stabilize NCR peptides during symbiosis.

host-microbe interactions | host peptides | signaling peptides | oxidative folding | regioisomers

Establishment of the ecologically, agriculturally, and economically important nitrogen-fixing symbiosis between legumes and rhizobia requires a mutual exchange of signals between the plant and its endosymbiotic bacterial partner (1). Development of the symbiosis is initiated when particular flavonoids secreted by the legume host induce the secretion of bacterial Nod factors. Nod factors activate signal transduction networks in the host plant, establish the nodule meristem necessary for nodule growth, stimulate the root hair curling that entraps rhizobia, and stimulate expression of an exopolysaccharide receptor. Synthesis of one or more rhizobial exopolysaccharides is then required for growth of infection threads, plant-derived tubes that permit the entrapped rhizobia to passage down the root hairs and penetrate into the interior of the developing nodule. In the infection zone of the nodule (the nondividing cells just after the nodule meristem), the bacteria are endocytosed into host membrane compartments, termed symbiosomes, where they undergo the major physiological changes necessary to enable nitrogen fixation (1).

Medicago truncatula and *Medicago sativa* (alfalfa) are inverted repeat-lacking clade (IRLC) legumes that form indeterminate nodules. After being endocytosed into the plant cytoplasm, their symbiotic partner *Sinorhizobium meliloti* undergoes terminal

differentiation into polyploid nitrogen-fixing bacteroids (2). It has been shown that one factor in promoting differentiation is the low levels of free oxygen in the plant cytoplasm due to leghemoglobin and the triggering of a hemoprotein sensor kinase, FixL, which starts a cascade to regulate *nif* gene transcription. A more recent advance in understanding the signaling events required for symbiosis has been the discovery that a family of >700 defensin-like nodule-specific cysteine-rich (NCR) peptides drive many of the complex physiological changes that characterize *S. meliloti* differentiation. These changes include alterations of the bacterial cell cycle to permit rounds of endoreduplication, major changes in patterns of transcription and translation, alterations of membrane permeability, and major morphological changes, including elongation and frequent branching (3–8). The NCR peptides are expressed only in nodules and are induced in successive waves as coordinated nodule development and rhizobial infection occur (5). The peptides are delivered to the symbiosome compartments through an endoplasmic reticulum-dependent secretion system that requires the action of the DNF1 nodule-specific signal peptidase (3, 4). The mature peptides are usually 20–55 aa long and vary greatly in their amino acid sequences; many can be clustered into two subgroups, NCR motif-A (four cysteines) and NCR motif-B (six cysteines) (9). Recent studies of two symbiotically defective *M. truncatula* mutants have shown that NCR169 (10) and NCR211 (11) are each required for a functional nitrogen-fixing symbiosis. Additionally, if *S. meliloti* carries a plasmid encoding the HrrP

Significance

Terminal differentiation of endocytosed rhizobia into nitrogen-fixing bacteroids inside of legume root nodules is orchestrated by a large family of host-encoded cysteine-rich signaling peptides. These peptides have diverse effects on bacteria, but their molecular mechanisms of action are still unknown. The presence of highly conserved cysteine motifs in this peptide family results in an additional layer of complexity because each individual peptide can potentially form several disulfide-cross-linked regioisomers under oxidative conditions. By demonstrating several distinct patterns of relative activities between the three disulfide-cross-linked regioisomers of NCR247, its reduced form, and a variant lacking cysteines, our work suggests that disulfide cross-linking can augment the rich complexity of *Rhizobium*-legume chemical communication required for symbiosis.

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(host range restriction peptidase) metalloprotease capable of degrading NCR peptides, symbiotic outcomes are affected at a late developmental stage (12). Although NCR peptides clearly play signaling roles in symbiosis, some members are bactericidal at higher concentrations. The membrane protein BacA plays a critical role in symbiosis by protecting *S. meliloti* cells in the symbiosome from the potential bactericidal action of NCR peptides (13, 14).

Prokaryotes, as well as lower and higher eukaryotes, use peptides as signaling molecules, and the cysteine-rich peptides (CRPs) constitute one of the most important families of these molecules. Particularly well-studied CRPs include conotoxins, neurotoxins produced by cone snails, and defensins, important effectors of innate immunity in vertebrates, invertebrates, insects, and plants (15). Although, *M. truncatula*'s >700 NCR peptides are distinct from its ~80 defensins (5), the shared structural feature of multiple disulfide bonds is thought to afford a compact structure and resistance to proteolytic and chemical degradation in both peptide families (16). Furthermore, reports indicate that reduced and oxidized forms of some CRP peptides have different biological activities (17–20). In principle, motif-A NCR peptides, which have four cysteine residues, can form three possible disulfide-cross-linked regioisomers whereas motif-B NCR peptides, which have six cysteine residues, can form 15 possible disulfide-cross-linked regioisomers. Nevertheless, there has been no systematic study of the relative effects of the various oxidized regioisomers and the reduced isoform on the symbiotic and bactericidal activities of any NCR peptide.

To address the issue of how NCR peptide structure and oxidation state relate to function, we chose the intensively studied NCR247 peptide (Fig. 1). NCR247 is composed of 24 aa, contains four cysteine residues, has a cationic charge (+5.6) at neutral pH, and also has the highest Boman index (4.6) of any of the NCR peptides (5). NCR247 is expressed in the older cells in the infection zone (zone II), where *S. meliloti* cells stop dividing and start differentiating, and also in interzone II and III, where a sudden growth of bacteroids is visible (21). Gene-expression profiling revealed that treatment of a synchronized *S. meliloti* population with a sublethal level of NCR247 affects ~15% of the genome, including three symbiotically important regulons (7). Two of these regulons are controlled by the ExoS/ChvI and FeuQ/FeuP two-component systems, respectively, and the other

by the master cell cycle regulator *ctrA*. Sublethal levels of NCR247 permit the cell to complete one round of DNA replication but then lead to inhibition of cell division, Z-ring formation, and septation (7). NCR247 also inhibits protein synthesis (21). Potential interacting partners of NCR247 by using affinity chromatography include FtsZ, numerous ribosomal proteins, the GroEL chaperone, and RNA polymerase units (21). Both the reduced isoform of NCR247 (NCR247_{red}) and one disulfide-linked NCR247 regioisomer have been shown to be bactericidal at higher concentrations (3, 22).

Our results demonstrate that the particular disulfide-cross-linked regioisomers of NCR peptides and the reduced NCR peptides can differ strikingly in their abilities to cause various biological effects. Taken together, these observations suggest that the NCR peptide-based language that plants use to control the development of their bacterial partner during symbiosis may be more complex than currently appreciated.

Results

Preparation of the Three Disulfide-Linked Regioisomers of NCR247.

Previous studies of the biological activities of NCR247 have used the reduced isoform NCR247_{red} (3, 22) or the disulfide-linked NCR247_{1-2,3-4} regioisomer (7, 14) (Fig. 1A). The latter form has been proposed (22) to be the isoform present in root nodules on the basis of Chou-Fasman predictions (14). To gain insights into the potential importance of disulfide bonds and cysteines to the biological activities of NCR247, we prepared the three disulfide-linked regioisomers of NCR247 (Fig. 1A). We then compared the biological activities of these oxidized peptides with those of NCR247_{red} and also of NSR247 (22). NSR247 is an NCR247 variant that cannot form disulfide bridges because all four cysteine residues are replaced by serines.

We obtained NCR247_{red} by Fmoc solid-phase peptide synthesis and used DMSO in oxidative folding assays. This mild oxidant has been used to fold basic and hydrophobic peptides, including human β -defensin 3 (23). We found that incubating NCR247_{red} in a 20% (vol/vol) DMSO solution (room temperature for 20 h) resulted in the formation of three species, which could be separated by HPLC (high-performance liquid chromatography) (Fig. 1B). Mass spectrometry revealed that each peak was a new species that had an m/z value 4 units less than that of NCR247_{red}. Thus, all three NCR247_{ox} regioisomers formed under these conditions, each of which contains two intramolecular disulfide bonds (Fig. 1C and D).

After HPLC separation and purification of each regioisomer (Figs. S1 and S2), we mapped the disulfide linkages of each peptide. We assigned the major peak (peak 2) to the NCR247_{1-4,2-3} regioisomer by identifying the peptide fragments generated by extended trypsin digestion (Fig. S3). To distinguish between NCR247_{1-2,3-4} and NCR247_{1-3,2-4}, which yield peptide fragments of identical mass after trypsin digestion, we synthesized the NCR247_{1-2,3-4} regioisomer by orthogonal protection of cysteines and subsequent pairwise deprotection and oxidation to use as an HPLC standard. Authentic NCR247_{1-2,3-4} displayed the same HPLC retention time as peak 3, thereby allowing us to infer that peak 1 is the NCR247_{1-3,2-4} regioisomer (Fig. S1). Our results suggest that NCR247_{1-4,2-3} is the thermodynamically most stable regioisomer of oxidized NCR247, rather than NCR247_{1-2,3-4} as had been proposed (14). It is possible that NCR247_{1-4,2-3} may be the most abundant isoform present in *M. truncatula* nodules, but other factors could influence the in vivo distribution of regioisomers. Circular dichroism (CD) spectroscopy did not reveal classical secondary structure features in any of the disulfide-cross-linked regioisomers (Fig. S4).

We also synthesized a derivative of NCR247_{red} in which fluorescein isothiocyanate was conjugated to the N terminus (FITC-NCR247_{red}). This derivative has cell division inhibition and bactericidal activity comparable with unmodified NCR247_{red} (Fig. S5). Because the robust optical signal provided by the fluorophore allows detection of relatively low concentrations of the peptide by HPLC, we used this peptide to evaluate whether NCR247_{red} stays reduced under our in vivo experimental conditions. After incubating

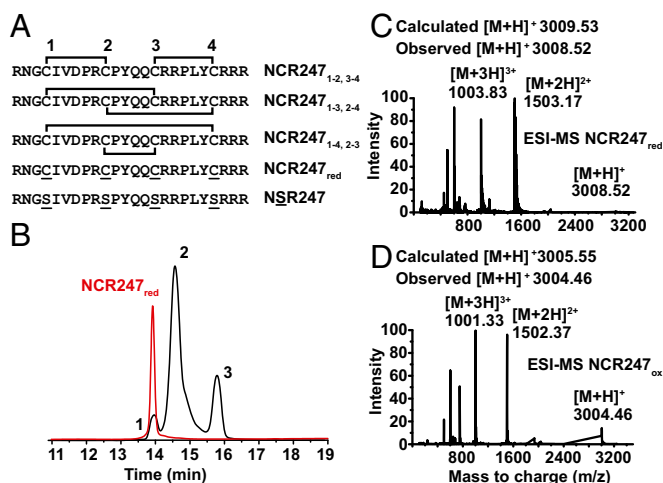


Fig. 1. Generation and analyses of NCR247 regioisomers. (A) Primary sequences and illustration of disulfide bond linkages in NCR247_{ox}. The brackets indicate the disulfide bond linkages in the oxidized peptides. (B) HPLC trace (10–60% acetonitrile over 30 min, 1 mL/min, 280 nm) showing that oxidation of NCR247_{red} in 20% DMSO generated three new peaks, which are assigned as the NCR247_{ox} regioisomers. (C and D) Deconvoluted ESI-MS spectra of NCR247_{red} and NCR247_{ox} variants indicate the oxidation state of the peptides.

a sublethal concentration (4 μM) of FITC-NCR247_{red} aerobically with *S. meliloti* cells for 3 h, the supernatant was analyzed by HPLC and mass spectrometry. The retention time and mass of the peptide recovered from the supernatant were the same as FITC-NCR247_{red} (Fig. S6). It is possible that some oxidation occurs within the bacterial cells; however, based on data presented below that indicate differences between oxidized and reduced forms in some bioassays, we reason that oxidation does not happen to a great extent.

We then systematically examined the relative abilities of sublethal doses of the three disulfide-linked regioisomers of NCR247, NCR247_{red}, and NSR247 to affect several physiological responses thought to be relevant to symbiosis: (i) the expression of representative ExoS/ChvI- and FeuQ/FeuP-controlled genes and the key regulatory gene *ctrA* (7), (ii) protein synthesis (21), and (iii) cell division, Z ring formation, and septum formation (3, 7). Additionally, we examined their relative bactericidal activities and resistance to proteolysis.

Effect of NCR247 Disulfide-Cross-Linked Regioisomers on ExoS/ChvI and FeuQ/FeuP Controlled Genes and *ctrA*. Our previous gene-expression profiling study of a synchronized population of *S. meliloti* treated with a sublethal concentration (4 μM) of NCR247_{1-2,3-4} (7) revealed the up-regulation of two symbiotically important regulons controlled by the ExoS/ChvI and FeuQ/FeuP two-component regulatory systems, respectively. The sensors, ExoS and FeuQ, are histidine kinases that phosphorylate their respective response regulators, ChvI and FeuP. The ExoS/ChvI regulon includes numerous genes required for the synthesis of the symbiotically important exopolysaccharide succinoglycan, as well as the known direct ChvI target *smc01581* (24). The FeuQ/FeuP regulon includes *ndvA*, which encodes an ABC cyclic glucan exporter required for symbiosis (25). In contrast to up-regulation of these two regulons, our transcriptional analysis showed that a sublethal concentration of the NCR peptide decreased the expression of *ctrA* (7), which encodes a critical cell cycle regulator and master regulator of symbiosis (26).

Using quantitative PCR (qPCR) to compare the expression levels of *smc01581* and *ndvA* genes after treatment of *S. meliloti* cultures with the NCR247 variants, we found that all disulfide-cross-linked regioisomers significantly up-regulated expression of both genes to similar extents. In contrast, NCR247_{red} had no effect, and NSR247 had only low activity in up-regulating *smc01581* and *ndvA* expression (Fig. 2A and B). We observed a different pattern of the relative bioactivities of the five NCR247 peptides when we examined their effects on *ctrA* expression (Fig. 2C). All three oxidized NCR247 regioisomers were again equivalently bioactive, reducing *ctrA* expression to similar degrees, and NSR247

was inactive. However, in contrast to having no effect on the expression of the ExoS/ChvI- and FeuQ/FeuP-regulated genes, NCR247_{red} was as active as the oxidized peptides in reducing *ctrA* expression. Taken together, these results reveal that the activity of the reduced isoform NCR247_{red} relative to that of the oxidized NCR247 regioisomers depends on the biological attribute being assayed.

NCR247 Regioisomers Selectively Inhibit Bacterial In Vitro Translation.

The protein translation machinery is one of the major targets of several signaling peptides in plant-microbe and insect-microbe symbioses (27). Farkas et al. (21) observed that the set of NCR247-interacting proteins identified by affinity chromatography includes ribosomal proteins and subsequently discovered that NCR247_{red} can inhibit protein synthesis in vitro (21). Consequently, we sought to determine whether the other NCR247 variants inhibit protein synthesis. Guided by the reported assay (21), we used an in vitro translation system in which translation of green fluorescent protein (GFP) mRNA can be monitored by measuring GFP fluorescence. We confirmed that NCR247_{red} (40 μM) substantially inhibits protein synthesis and observed that the relative abilities of the NCR247 variants to inhibit translation were strikingly different from those we had observed in our transcription assays (Fig. 2D). NCR247_{1-2,3-4} and NCR247_{1-3,2,4} inhibited translation, albeit to a lesser degree than NCR247_{red}. In striking contrast, NCR247_{1-4,2-3} was highly effective at inhibiting the translation of GFP (Fig. 2D) and could even interfere with translation at the sublethal level of 4 μM (Fig. S7). Moreover, NSR247 was less effective at inhibiting translation compared to NCR247_{red}, which indicates that the cysteine residues are generally important but not essential for this activity (Fig. 2D).

NCR247 Cell Division Inhibition Activity Is Independent of Disulfide Connectivity, and Cysteines Are Not Required.

We have previously shown that a sublethal concentration (4 μM) of NCR247_{1-2,3-4} does not alter the initiation or progression of DNA replication, such that the cellular DNA content increases from 1C to 2C, for synchronized *S. meliloti* cultures at the G1 phase. However, NCR247_{1-2,3-4} specifically and robustly blocks cell division in *S. meliloti* in G2 (7). At the sublethal dose of 4 μM , the three NCR247_{ox} regioisomers, NCR247_{red}, and NSR247 all exhibited similar abilities to prevent the 2C cells from dividing to yield daughter cells with 1C DNA content (Fig. 2E). Lowering the peptide concentration to 2 μM revealed that one of the minor regioisomers, NCR247_{1-2,3-4}, was somewhat more active than the other peptides (Fig. 2F and Fig. S8). We have previously shown that NCR247_{1-2,3-4} causes an early block in cell division because very little septation is observed and Z-ring formation is inhibited

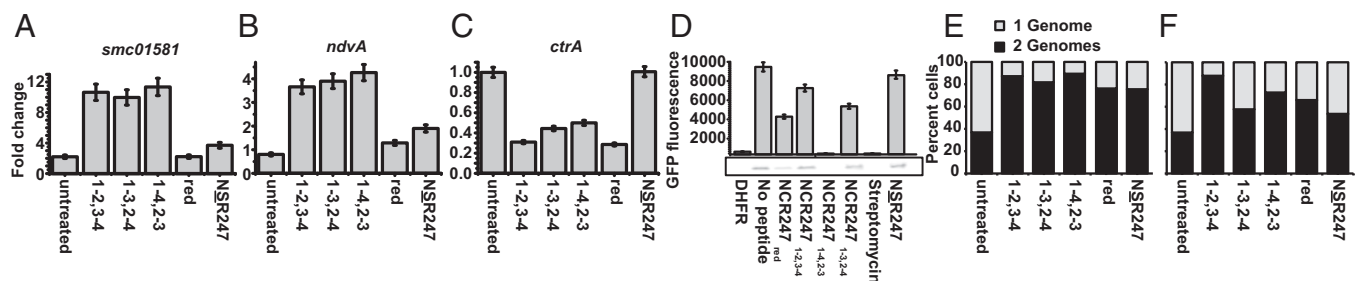


Fig. 2. Treatment with NCR247 variants affects expression levels of genes regulated by the ExoS-ChvI, FeuQ-FeuP regulons and of the master cell-cycle regulator *ctrA*. (A and B) Fold change in *smc01581* (A) and *ndvA* (B) expression relative to *smc00128* after 15 min with no peptide or with a sublethal concentration (4 μM) of peptide-treated synchronized cultures. (C) The *ctrA* expression in nontreated and peptide-treated synchronized cultures relative to *smc00128*. (D) NCR247_{red} and NCR247_{1-4,2-3} specifically inhibit protein synthesis in vitro. (D, Upper) NCR247 variants (40 μM) were added to the "in vitro translation reaction" mixture, and GFP fluorescence was monitored by using a fluorescence plate reader after 3 h ($\lambda_{\text{ex}} = 485 \text{ nm}$, $\lambda_{\text{em}} = 530 \text{ nm}$). (D, Lower) In-gel fluorescence of GFP reporter protein after 3 h of treatment with peptide. (E and F) Level of cell division inhibition activity of NCR247 is affected by redox state and oxidative folding. (E and F) DNA content per cell during synchronized growth of *S. meliloti* cultures treated with or without 4 μM (E) and 2 μM (F) of NCR247 variants. DNA content per cell during synchronized growth of *S. meliloti* cultures treated with or without NCR247 variants. Flow cytometry was used to measure the percentage of cells with one genome or two genomes. All error bars represent SD from the mean, based on triplicate experiments.

(7). The other two NCR247_{ox} regioisomers, NCR247_{red}, and NSR247, also inhibited the appearance of septa to a similar degree (Fig. S9A). Moreover, when we used an inducible FtsZ-GFP translational fusion protein to visualize the Z-ring in cells in synchronous cultures at the G2 phase (7), we found that each of the five NCR247 variants (4 μ M) exhibited similar abilities to inhibit Z-ring formation (Fig. S9B).

Taken together, these results show that the pattern of activities of the various NCR peptides in inhibiting cell division differs from the activity patterns obtained from the transcription and translation bioassays. Even though NCR247_{1-2,3-4} was the most active at 2 μ M, the fact that the three disulfide-cross-linked regioisomers had approximately the same activity as NCR247_{red} at 4 μ M indicates that disulfide bond formation is not crucial. Our observation that the NSR247 peptide is active further indicates that none of the cysteines are required for inhibition of cell division, thereby suggesting a different underlying molecular mechanism from those involved in regulation of transcription and translation.

Disulfide Cross-Linking Influences the Relative Bactericidal Activities of NCR247 Peptides in Wild-Type *S. meliloti*. Although both NCR247_{red} and NCR247_{1-2,3-4} induce bacteroid-like features in free-living *S. meliloti* at sublethal concentrations (3, 7, 14), both peptides reduce the colony-forming ability of *S. meliloti* at higher concentrations (14, 22). We compared the bactericidal activity of the three NCR247_{ox} regioisomers (Fig. 3A) and observed that NCR247_{1-4,2-3} and NCR247_{1-2,3-4} are both ≥ 10 -fold less active than NCR247_{red} and NCR247_{1-3,2-4}. In this respect, our results for NCR247_{1-3,2-4} differ from those of Haag et al. (22), who reported that NCR247_{1-3,2-4} is less bactericidal than NCR247_{1-2,3-4} (22). However, they used a preparation of NCR247_{1-3,2-4} that was known to be impure (22), and we now have shown that the impurities in a sample of their peptide are not other NCR247_{ox} regioisomers as they had speculated (22) (Fig. S10). Thus, our data indicate that disulfide bonds are not required for NCR247 to be bactericidal against *S. meliloti* although they can modulate its activity. Of the oxidized species, the most abundant regioisomer, NCR247_{1-4,2-3}, has the weakest bactericidal activity against *S. meliloti*, an observation of potential physiological significance. NSR247 has negligible bactericidal activity (Fig. 3A), which indicates that one or more cysteines residues are important for the bactericidal activity of NCR247 against *S. meliloti* even though disulfide bonds do not seem to be required.

BacA Function Attenuates the Bactericidal Activity of All NCR247 Regioisomers and NCR247_{red} Against *S. meliloti*. The membrane protein BacA is required for *S. meliloti* to establish the chronic intracellular infection that underlies its symbiosis (13) and also for the related mammalian pathogen *Brucella abortus* to establish the chronic intracellular infection that underlies its pathogenesis (28). We found that an *S. meliloti* Δ *bacA* mutant also displayed increased sensitivity to NCR247_{1-3,2-4} and NCR247_{1-4,2-3}, as well as

NCR247_{red}. Thus, our results indicate that the molecular mechanism for BacA-mediated resistance of *S. meliloti* to bactericidal NCR247 peptide works equally well for all conformations of the NCR247 peptide.

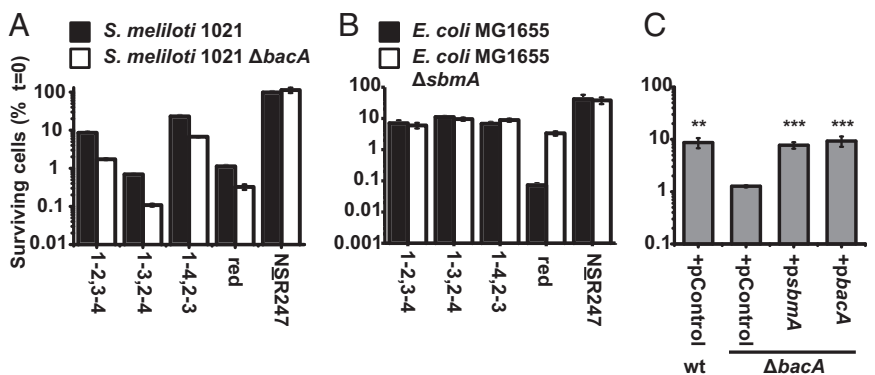
As shown for NCR247_{1-2,3-4}, the sensitivity of the Δ *bacA* mutant to killing by all of the NCR247 peptides can be restored to that of a WT strain by expressing a plasmid-borne copy of the *S. meliloti* *bacA*⁺ gene or its isofunctional *Escherichia coli* ortholog, *sbmA*⁺ gene (Fig. 3C). Interestingly, despite *E. coli* *SbmA* being isofunctional with BacA when expressed in *S. meliloti* (29), we observed that an *E. coli* Δ *sbmA* mutant did not display increased sensitivity to killing by any of the NCR247_{ox} regioisomers and was more resistant, rather than more sensitive, to NCR247_{red} (Fig. 3B). *E. coli* Δ *sbmA* mutants have previously been shown to exhibit resistance to killing by several proline-rich peptides. The mechanism of *S. meliloti*'s BacA-mediated resistance to NCR247 will require further investigation and may be complex.

Indeed, BacA/SbmA has been shown to affect both the modification of *S. meliloti* and *B. abortus* lipid A by very long chain fatty acids (VLCFAs) and import of proline-rich peptides. Furthermore, the differing responses of the *S. meliloti* *bacA* and *E. coli* *sbmA* mutants to NCR247 peptides indicate that the physiological role of the conserved BacA/SbmA protein in a particular bacterium can also be influenced by other cellular functions. As symbiosis proceeds, the expression of NCR247 and two other intensively studied NCR peptides (NCR211 and NCR169) roughly coincides with *bacA*, consistent with BacA protecting the nascent bacteroids from the bactericidal property of the NCR247 peptide. The expression of genes encoding functions that could influence the properties and functions of NCR peptides, such as bacterial periplasmic disulfide, oxidoreductases, and glutathione-S-transferases, also varies during the development of the symbiosis (30) (Fig. S11).

Disulfide Bonding in NCR247 Protects Against Degradation by HrrP.

Although NCR peptides have been proposed to be oxidized in planta (9), our results confirm that NCR247_{red} is active in several symbiotically related bioassays. A possible role for disulfide cross-linking of NCR peptides could be to stabilize them against proteolytic degradation (31). Expression of the plasmid-encoded metalloprotease HrrP, which is capable of degrading a range of NCR peptides in vitro, has recently been shown to restrict nodule development and nitrogen fixation of certain plant hosts (12). To examine whether disulfide bond formation modulates the proteolytic stability of NCR247, we performed degradation assays using HrrP and catalytically inactive variant HrrP (E62A). The assays were done using the oxidized and reduced NCR variants as substrates. All three NCR247_{ox} regioisomers were markedly less susceptible to degradation by HrrP than NCR247_{red} (Fig. 4). At a time point when NCR247_{red} was 80% degraded, NCR247_{1-3,2-4} and NCR247_{1-4,2-3} exhibited virtually no degradation, and NCR247_{1-2,3-4}

Fig. 3. Antimicrobial activity of NCR247 depends on the regioisomer. (A) *S. meliloti* Sm1021 WT and Δ *bacA* mutant strains were treated with 20 μ M NCR247 variants for 180 min in MOPS-GS buffer supplemented with 1% casamino acids. (B) *E. coli* MG1655 WT and Δ *sbmA* mutant strains were treated with 6 μ M NCR247 variants for 180 min in 10 mM sodium phosphate buffer (pH 7.0). (C) Bars show the relative survival of the defined *S. meliloti* strains with either empty control plasmid pRK404, plasmid pA1351 harboring the *E. coli* *sbmA* gene, or plasmid pJG51A harboring the *S. meliloti* *bacA* gene. Cells were treated with 20 μ M NCR247_{1-2,3-4} for 180 min in MOPS-GS buffer supplemented with 1% casamino acids. Bars show bacterial survival assessed relative to the cfu·mL⁻¹ at t = 0 min. All error bars represent mean \pm SD (n = 3) based on three technical replicates representing trends observed in at least two independent experiments. Significant values are **P \leq 0.01 and ***P \leq 0.001.



susceptible bacterium that manages to coinfect along with the rhizobial symbiont. Along these lines, the BacA protein plays a critical role in symbiosis by preventing the rhizobia from being killed by the concentrations of NCR peptides present in the symbiosome, so that the signaling roles of the NCR peptides can predominate.

NCR peptides affect multiple components of bacterial physiology to achieve a successful symbiosis, some inducing the final terminal differentiated state and others affecting bacteroid persistence. Some NCR peptides may work cooperatively with other NCR peptides to exert positive or negative effects on affected pathways whereas multiple NCR peptides may bind to the same target to achieve their ultimate biological effects. Although some NCR peptides are likely to be at least partially functionally redundant with other members of the large NCR peptide family, at least two have unique functions essential for symbiosis (NCR169 and NCR211) (10, 11). Which physiological functions require peptide interactions with membranes and which may require interactions with specific cellular proteins remain to be clarified. Cationic peptides can disrupt cellular membranes, consequently some cationic NCR peptides may affect membrane permeability, leading to physiological outcomes and also permitting other classes of NCR peptides to enter the bacteria. Interactions of an NCR peptide with a specific protein target would probably be strongly affected by the pattern of disulfide cross-linking.

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Materials and Methods

Detailed materials and methods can be found in *SI Materials and Methods*.

Bacterial Growth Conditions. For all experiments, *S. meliloti* 1021 WT and Δ bacA-null mutant strains were grown in Lysogeny broth (LB) prepared with 5 g·L⁻¹ NaCl and supplemented with 2.5 mM CaCl₂ and 2.5 mM MgSO₄ (LB-MC) for 48 h at 30 °C. *E. coli* WT and Δ sbmA-null mutant strains were grown in LB (5 g·L⁻¹ NaCl) at 37 °C. Unless indicated otherwise, *S. meliloti* cultures were supplemented with 200 μ g·mL⁻¹ streptomycin (Sm).

Folding and Disulfide Formation. The folding reaction typically proceeded at room temperature overnight in a sealed vial with gentle stirring. Oxidation of NCR247_{red} was done in 20% DMSO in water (23, 37).

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