

# Recombinant Macrocyclic Lanthipeptides Incorporating Non-Canonical Amino Acids

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## S Supporting Information

**ABSTRACT:** Nisin is a complex lanthipeptide that has broad spectrum antibacterial activity. In efforts to broaden the structural diversity of this ribosomally synthesized lantibiotic, we now report the recombinant expression of Nisin variants that incorporate noncanonical amino acids (ncAAs) at discrete positions. This is achieved by expressing the *nisA* structural gene, cyclase (*nisC*) and dehydratase (*nisB*), together with an orthogonal nonsense suppressor tRNA/aminoacyl-tRNA synthetase pair in *Escherichia coli*. A number of ncAAs with novel chemical reactivity were genetically incorporated into NisA, including an  $\alpha$ -chloroacetamide-containing ncAA that allowed for the expression of Nisin variants with novel macrocyclic topologies. This methodology should allow for the exploration of lanthipeptide variants with new or enhanced activities.

The lanthipeptides are a subclass of a large family of ribosomally synthesized and post-translationally modified macrocyclic peptides (RiPPs).<sup>1</sup> Members that possess antibacterial activity are known as lantibiotics,<sup>2</sup> and are defined by characteristic lanthionine or methyllanthionine thioether bridges that impart backbone rigidity. The most studied member of the lanthipeptides, Nisin A, depicted as its prepeptide or mNisA (Figure 1), has been used as a food preservative for the past 50 years.<sup>3</sup> It features multiple dehydrated amino acids arising from the activity of a dehydratase (NisB) on serine (Ser) and threonine (Thr) residues, and five thioether cross-links (rings A, B, C and D/E) formed by a cyclase (NisC) that catalyzes thia-Michael reactions.<sup>4</sup> Nisin exerts its antimicrobial activity through its N-terminal domain that recognizes and sequesters the pyrophosphate moiety of Lipid II, and the C-terminal domain that inserts into the outer membrane.<sup>5,6</sup> This dual mode of action, directed toward nonproteinaceous bacterial targets, has slowed the development of resistant strains.<sup>7</sup> However, its poor aqueous solubility and pH sensitivity have delayed its use as a human therapeutic.<sup>8</sup>

To further explore the potential utility of this complex natural product, various efforts have been undertaken to generate variants of Nisin and other lantibiotics,<sup>9</sup> including the use of solid phase synthesis,<sup>10–12</sup> biomimetic approaches,<sup>13</sup> the in vitro biosynthesis of lanthionine-containing peptides<sup>14,15</sup> and an in vitro mutasynthetic approach.<sup>16</sup> To expand the number of amino acids building blocks that can be biosynthetically incorporated into lanthipeptides, we now report that orthogonal nonsense suppressor tRNA/aminoacyl-tRNA synthetase pairs

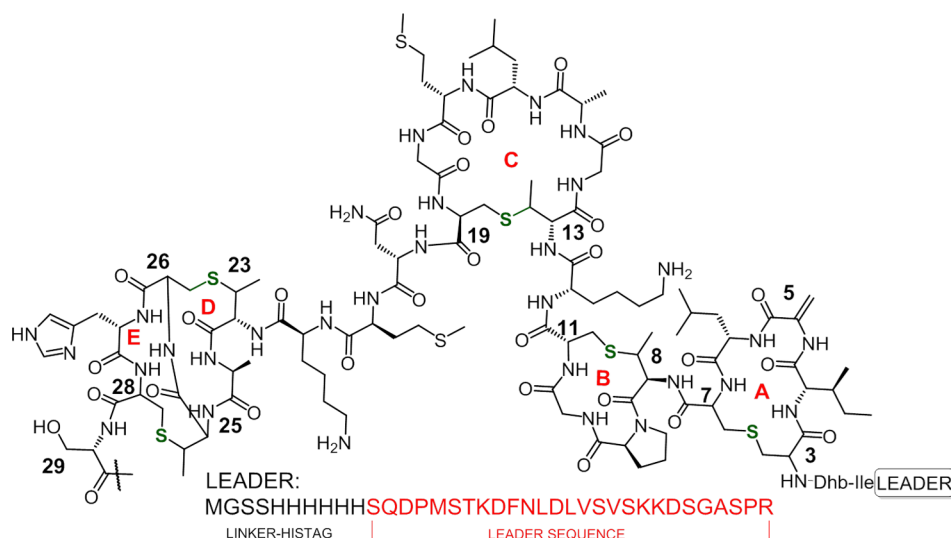
(tRNA/aaRS) can be used to site-specifically insert ncAAs into Nisin in bacteria.<sup>17</sup>

To date, two examples of ncAA incorporation into lanthipeptides using nonsense codon suppression have been reported by van der Donk.<sup>18,19</sup> Kuipers also reported the incorporation of tryptophan analogues into Nisin using an auxotrophic strain.<sup>20</sup> With the goal of creating Nisin variants through biomimetic cyclization reactions (Figure 2), we built upon the former system<sup>18</sup> and began by recombinantly expressing wild-type (WT) NisA in bacterial strains in which we have previously genetically encoded a large number of ncAAs. The *nisA* structural gene together with the dehydratase (*nisB*) were inserted into a pRSF-1 vector to afford pRSF-NisA. The cyclase, *nisC*, was inserted into a pACYC backbone to afford pACYC-NisC. The Ser codon at position 5 of *nisA* was then mutated to the amber stop codon TAG (Ser5TAG), and the resulting plasmid (pZC1) was coexpressed, along with pACYC-NisC, and a pULTRA plasmid harboring either a polyspecific *Methanocaldococcus jannaschii* tRNA<sup>Tyr</sup>/TyrRS or *Methanosarcina barkeri* pyrrolysine tRNA<sup>Pyl</sup>/PylRS.<sup>21</sup> We then attempted to substitute a number of ncAAs containing diverse side chains including keto, azide and acetylene groups for Ser5 (Figure 3A). SDS-PAGE and MS analysis (Figure S1–S3) confirmed incorporation of pAcF, pAzF and ProCk at position 5 in yields ranging from 3 to 5 mg/L (Figure 3B). Although promising, in our hands the expression system afforded incomplete dehydration of Ser and Thr residues, giving rise to a distribution of products.

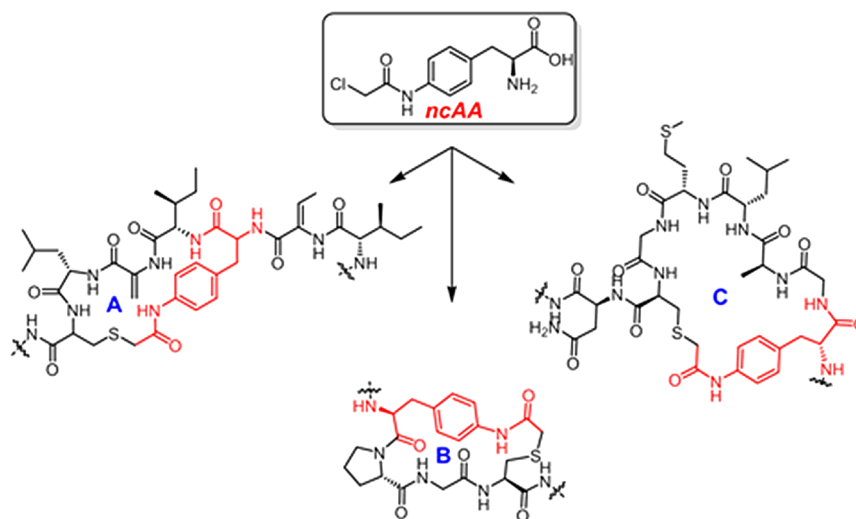
To solve this issue, we capitalized on the recent finding that the dehydratase NisB requires glutamylated tRNA (tRNA<sup>Glu</sup>) in order to acylate the Ser/Thr side chains of NisA with glutamate prior to dehydration.<sup>22</sup> We reasoned that overexpressing tRNA<sup>Glu</sup> could improve the dehydration efficiency. To test this notion, we constructed a plasmid (pZC2) bearing *Escherichia coli* tRNA<sup>Glu</sup>/GluRS downstream of a proK constitutive promoter encoded on pACYC-NisC. Coexpression of pZC2 with pRSF-NisA WT, showed a substantial increase in dehydration efficiency (WT mNisA = 7299 Da), confirming our hypothesis (Figure S4, S5).

Given this optimized system, we next explored the possibility of taking advantage of electrophilic ncAAs and the innate reactivity of cysteine residues present in NisA, to generate Nisin variants with altered ring structures. We began by incorporating mildly reactive ncAAs<sup>23</sup> in place of Ser/Thr residues at lanthionine ring junctions, starting with the ring A Ser3TAG

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**Figure 1.** Wild-type mNisA. Highlighted thioether rings and key residues for this study (residues 30–34 omitted for simplicity).



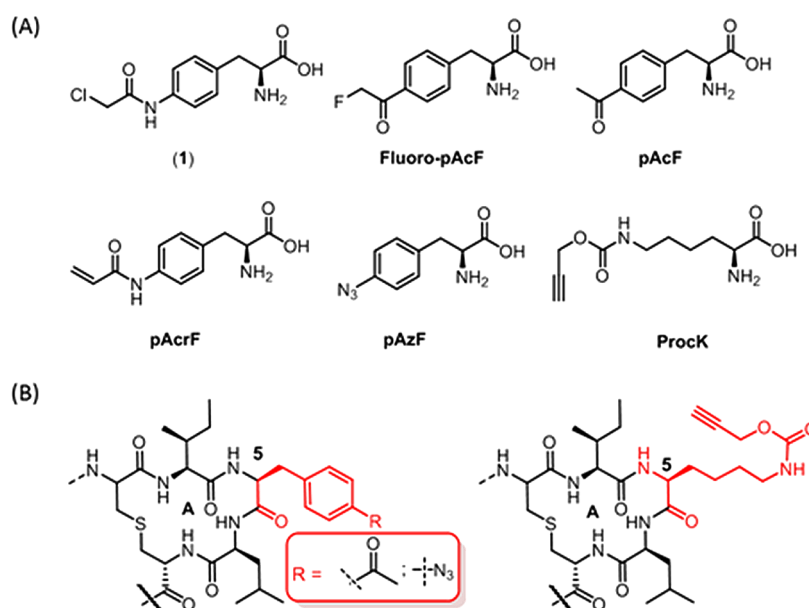
**Figure 2.** Altered ring structures of Nisin created by ncAA incorporation.

mutant (pZC3). Fluoro-pAcF (Figure 3A) was incorporated into NisA at residue 3 in a yield of 2 mg/L. MS analysis showed a mass (7418 Da) corresponding to the loss of fluoride (Figure S6). This result, although encouraging, was somewhat cryptic due to the similar molecular weight of fluorine and water. Because water acts as a leaving group during precursor peptide maturation, an extra post-translational dehydration would result in a product with a similar mass as that of the desired, cyclized peptide. This led us to reconsider the ncAA to be used.

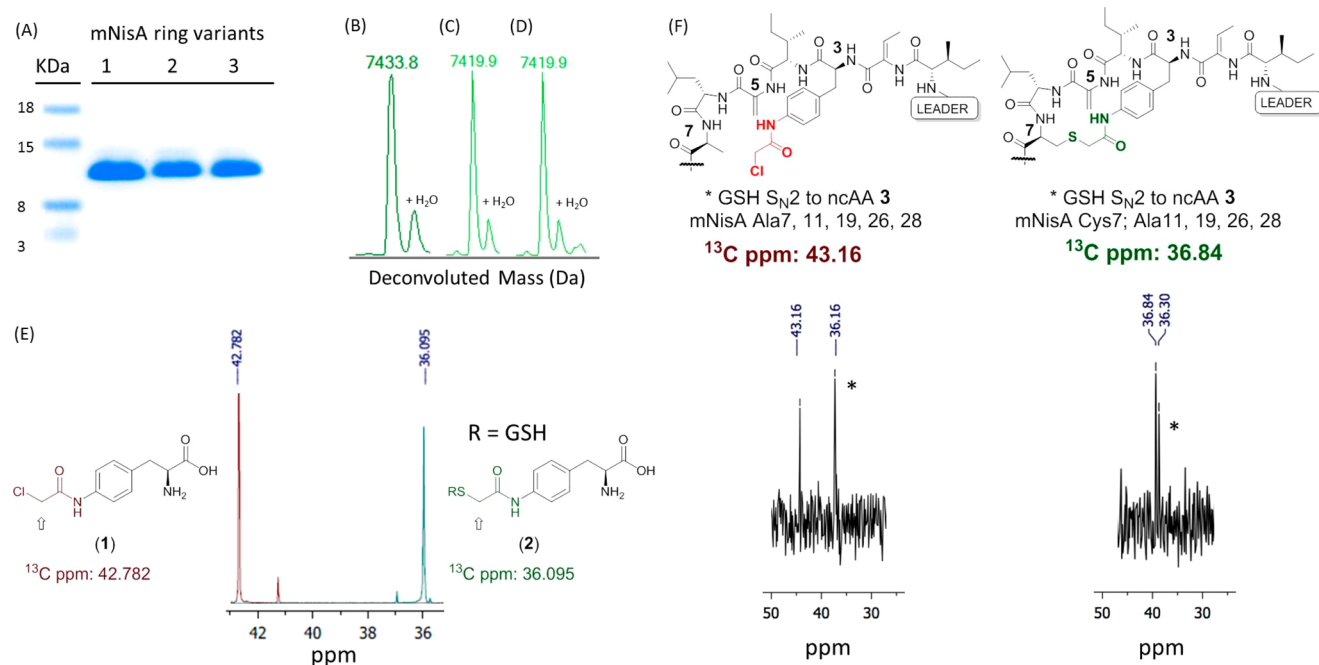
We reasoned that a chloroacetamide ncAA<sup>24</sup> (Figure 3A, 1) would be the best compromise in terms of reactivity and leaving group MW for mass analysis. Gratifyingly, when 1 was incorporated into ring A of mNisA (Ser3TAG), MS analysis showed a mass of 7433 Da  $[M/z]^{+1}$ , consistent with a fully dehydrated and cyclized peptide containing a novel ring A thioether architecture. We next investigated whether this approach could be extended to other ring junctions as a general strategy toward lanthipeptides diversification. To this end, the remaining dehydratable residues (Ser/Thr) involved in ring cyclization were individually point mutated to TAG and the resulting constructs cotransformed with pULTRA and pZC2. Upon expression and isolation of the prepeptides, SDS-

PAGE (Figure 4A) and mass analysis revealed cyclization of thioether rings A, B and C, corresponding to the Ser3TAG, Thr8TAG and Thr13TAG mutations, respectively, in yields of 1.5 to 3 mg/L (Figure 4B, C and D, Figure S7–S9). Efforts to create intertwined thioether ring D/E (Thr23TAG and Thr25TAG) were troublesome, giving rise to a mixture of desired and truncated products at Lys22.

Our next goal was to further characterize the newly formed bond in the Ser3 to 1 mutant. To simplify analysis, we decided to generate a monocyclic Nisin. To this end, we constructed two plasmids derived from the common vector, NisA Ser3TAG (pZC3). In the first construct (pZC4), we sequentially mutated all cysteine residues to Ala (Cys7, 11, 19, 26, 28Ala). The second construct (pZC5) retained Cys7 and all the remaining Cys were mutated to Ala (Cys11, 19, 26, 28Ala). Each construct was individually cotransformed with the accessory plasmids (pULTRA, pZC2), expressed in the presence of 1, and the mature peptides were isolated. MS analysis confirmed cyclization only for the peptide bearing reactive Cys7 (Figure S11). In contrast, the fully mutated Cys to Ala construct showed a MS spectrum for a peptide carrying the unreacted chloroacetamide. Interestingly, this noncyclized peptide showed



**Figure 3.** (A) ncAAs used in this study. (B) Biorthogonal chemical handles incorporated into SerSTAG NisA.



**Figure 4.** (A) SDS-PAGE mNisA ring variants incorporating **1**. Lane 1, ring A; lane 2, ring B; lane 3, ring C. (B–D) QTOF-ESI for mNisA ring variants incorporating **1**: (B) Ser3TAG, ring A. (C) Thr8TAG, ring B. (D) Thr13TAG, ring C. (E)  $^{13}\text{C}$  NMR of labeled **1** (red) and **2** (green). (F)  $^{13}\text{C}$  NMR mNisA Cys to Ala mutants incorporating labeled **1**, noncyclized (red) and cyclized (green) peptides. Asterisks indicate GSH addition products.

an extra dehydration event that was ablated upon Ser29 to Ala mutation in the core peptide (Figure S10). This observation is in accordance with the hypothesis that dehydration and cyclization events in Nisin biosynthesis are coupled, and that precise PTM control depends upon conformationally restricted substrates.<sup>25,26</sup>

To obtain NMR evidence for the novel thioether linkage, we synthesized isotopically labeled ncAA (**1**) bearing a  $^{13}\text{C}$  at the reaction center (2-chloroacetamide position) using chloroacetyl chloride-2- $^{13}\text{C}$  as a starting material. Incorporation of  $^{13}\text{C}$ -**1** into Nisin mutants should afford chemical shift differences

between cyclized and noncyclized product. As a control experiment, we recorded the  $^{13}\text{C}$  NMR spectrum of labeled **1**, and then treated it with excess glutathione (GSH) to form a thioether bond as a standard. A substantial chemical shift difference between the starting ncAA-chloroacetamide (42.78 ppm) and the GSH-displaced product **2** (36.09 ppm) was observed (Figure 4E). We then expressed and HPLC purified mutants Ala7, Ala11, Ala19, Ala26, Ala28 and Cys7, Ala11, Ala19, Ala26, Ala28 encoded on pZC4 and pZC5, respectively, incorporating  $^{13}\text{C}$ -(**1**). The peptide lacking all cysteines showed  $^{13}\text{C}$  NMR peaks consistent with the presence of a

chloroacetamide (43.16 ppm) and its GSH adduct. On the other hand, the peptide harboring reactive Cys7, showed a  $^{13}\text{C}$  resonance (36.84 ppm) consistent with the standard thioether **2**, confirming the macrocyclization event (Figure 4F). The additional thioether resonance (36.30 ppm) is likely due either to GSH addition to **1**, with Cys7 cyclizing to dehydroalanine 5 (Dha5), or GSH addition to Dha5 of the expanded cyclic peptide. Upon trypsin-mediated cleavage of the leader peptide, Nisin analogues bearing novel thioether linkages were tested for antibacterial activity. Unfortunately, although a halo assay against *Micrococcus luteus* indicated that this Nisin ring variants are devoid of antibacterial activity (whereas WT Nisin retained activity) (Figure S19), the possibility of constructing non-natural lanthipeptides with positional precision should allow for the interrogation of broader chemical space.

In summary, we genetically incorporated diverse ncAAs containing either biorthogonal handles or mildly reactive functional groups into Nisin A. NMR and tandem MS spectroscopy confirmed the formation of a novel ring A thioether bond for one of the mutants. We are currently probing various ring topologies with the aim to restore biologically active conformations. The ability to diversify lanthipeptide macrocycles at the ribosomal stage may lead to improvements in their pharmacological properties.

## ■ ASSOCIATED CONTENT

### Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/jacs.7b04159.

Mass spectra (ESI-QTOF, LC MS/MS), NMR data, cloning and synthetic protocols (PDF)

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### Notes

The authors declare no competing financial interest.

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