Increased D-alanylation of lipoteichoic acid and a thickened septum are main determinants in the nisin resistance mechanism of *Lactococcus lactis*

Naomi E. Kramer,¹,² Hester E. Hasper,² Patrick T. C. van den Bogaard,³ Siegfried Morath,⁴ Ben de Kruijff,² Thomas Hartung,⁴ Eddy J. Smid,⁵ Eefjan Breukink,² Jan Kok¹ and Oscar P. Kuipers¹

Correspondence
Oscar P. Kuipers
o.p.kuipers@rug.nl

¹Molecular Genetics Group, Groningen Biomolecular Sciences and Biotechnology Institute, University of Groningen, PO Box 14, 9750 AA Haren, The Netherlands
²Department of Biochemistry of Membranes, Center for Biomembranes and Lipid Enzymology, Institute of Biomembranes, Utrecht University, Padualaan 8, 3584 CH Utrecht, The Netherlands
³Eukaryotic Microbiology, Groningen Biomolecular Sciences and Biotechnology Institute, University of Groningen, PO Box 14, 9750 AA Haren, The Netherlands
⁴European Commission, Joint Research Centre, IRCP, European Centre for the Validation of Alternative Methods, 21020 Ispra, Italy
⁵NIZO Food Research, Flavour and Natural Ingredients Section, PO Box 20, 6710 BA Ede, The Netherlands

Received 27 November 2007
Revised 1 February 2008
Accepted 22 February 2008

Nisin is a post-translationally modified antimicrobial peptide produced by *Lactococcus lactis* which binds to lipid II in the membrane to form pores and inhibit cell-wall synthesis. A nisin-resistant (NisR) strain of *L. lactis*, which is able to grow at a 75-fold higher nisin concentration than its parent strain, was investigated with respect to changes in the cell wall. Direct binding studies demonstrated that less nisin was able to bind to lipid II in the membranes of *L. lactis* NisR than in the parent strain. In contrast to vancomycin binding, which showed ring-like binding, nisin was observed to bind in patches close to cell-division sites in both the wild-type and the NisR strains. Comparison of modifications in lipoteichoic acid of the *L. lactis* strains revealed an increase in D-alanyl esters and galactose as substituents in *L. lactis* NisR, resulting in a less negatively charged cell wall. Moreover, the cell wall displays significantly increased thickness at the septum. These results indicate that shielding the membrane and thus the lipid II molecule, thereby decreasing abduction of lipid II and subsequent pore-formation, is a major defence mechanism of *L. lactis* against nisin.

**INTRODUCTION**

*Lactococcus lactis* is a lactic acid bacterium (LAB) that is of major importance in the food industry. Some LAB produce agents that are capable of inhibiting the growth of a wide variety of food spoilage bacteria, for example by the excretion of antimicrobial peptides or bacteriocins. One of the best-studied antimicrobials is nisin (Sahl & Bierbaum, 1998). Nisin, a positively charged lantibiotic produced by *L. lactis*, and nisin-producing strains have been used as food preservatives for a long time, especially in dairy products (Delvès-Broughton et al., 1996; Hurst, 1981; Mattick & Hirsch, 1944). Nisin displays a broad spectrum of activity against Gram-positive bacteria (Jung, 1991) and kills bacteria by inhibiting cell-wall biosynthesis and by forming pores in the cytoplasmic membrane, both initiated by binding to lipid II (Breukink et al., 1999; Brötz et al., 1998). Multiple molecules in the lipid II–nisin complex aggregate to form a pore (Hasper et al., 2004). Recently, it has been reported that nisin, as well as some non-pore-forming lantibiotics are also able to abduct lipid II from major sites of cell-wall biosynthesis, providing another mechanism of cell killing (Hasper et al., 2006). Insights into the molecular mechanisms by which bacteria develop nisin resistance and the basis of the large differences in nisin sensitivity observed among wild-type bacterial species is of major importance for future antimicrobial applications of nisin or nisin variants.

Abbreviations: LTA, lipoteichoic acid; WTA, wall teichoic acid.
NMR spectra are available with the online version of this paper.
sensitivity are not correlated, but that resistance is a combined result of changes in the cell wall that hamper nisin from reaching lipid II (Kramer et al., 2004). The cell wall in Gram-positive bacteria consists of a relatively thick, multi-layered peptidoglycan sacculus that, depending on the bacterial species, may contain proteins, lipoteichoic acid (LTA), wall teichoic acid (WTA) and polysaccharides. Notably, not all Gram-positive bacteria harbour LTA or WTA. The cell wall has a net negative charge, mainly because of the LTA and WTA content. LTA in its most common form consists of a polyglycerol phosphate that is linked to the membrane via a glycolipid anchor. Alternatively, a polyribitol phosphate may form the LTA core, but other polyols such as mannotol, erythritol or arabitol can be present (Naumova et al., 2001). Some bacteria do not possess LTA, but have other anionic polymers such as teichuronic acid. LTA can contain various glycosyl substituents, such as an OH group, D-glucosamine and D-galactose (Delcour et al., 1999). In Gram-positive bacteria, four proteins, encoded by the dlt operon, are required for the synthesis of D-alanyl esters for substitution in LTA, which results in positive charges being incorporated into the mostly negatively charged cell wall (Delcour et al., 1999; Peschel et al., 1999). To date, it is not known whether or not L. lactis contains WTA. Three functions of D-alanyl teichoic acids have been proposed so far, namely (i) preventing the binding of autolysins (Steen et al., 2003), (ii) maintaining cation homeostasis, and (iii) defining the electrochemical properties of the cell wall (Heaton & Neuhaus, 1993).

LTA has been shown to play an important role in nisin resistance in Staphylococcus aureus and Streptococcus bovis; however, the exact LTA composition of these strains has not been analysed. A Staphylococcus aureus strain containing several copies of the dlt operon is less sensitive to various antimicrobial peptides, including nisin (Peschel et al., 1999). Nisin-resistant (NisR) cells of Streptococcus bovis harbour more LTA than nisin-sensitive cells (Mantovani & Russell, 2001). LTA alanylation also seems to affect the susceptibility of Streptococcus pneumoniae to nisin (Kovacs et al., 2006). We have shown previously that the cell wall of Gram-positive bacteria can be a barrier for nisin to reach lipid II, since protoplasts of a NisR strain of Micrococcus flavus were almost as sensitive to nisin as protoplasts of the parent strain (Kramer et al., 2004). However, the LTA of the NisR strains has not been isolated in any of these studies, nor have the molecular changes in LTA been reported to be involved in the nisin resistance mechanism.

A transcriptome analysis study has shown that the expression of various genes involved in four possible mechanisms was changed in a L. lactis NisR strain compared to the wild-type (Kramer et al., 2006), based on the following observations. (i) Genes involved in cell-wall structure showed higher expression of the dlt and gal operons, resulting in a changed LTA structure. GalE, encoded by the gal operon in L. lactis subsp. cremoris (Grossiord et al., 2003), is involved in galactose substitution in LTA. pbp2A was also more highly expressed. There was also (ii) a difference in the expression of ABC transporters, (iii) a difference in the expression of genes involved in the saturation and elongation of fatty acids, and (iv) changes in the expression of genes possibly controlling the pH on the outer side of the membrane, preventing nisin binding to lipid II (Kramer et al., 2006). However, the biochemical changes in membrane and cell wall were not further investigated in this study.

Since earlier studies have indicated that several genes involved in cell-wall synthesis contribute to nisin resistance (Kramer et al., 2006), and do not influence the total lipid II content, we set out to further investigate the role of the cell wall in nisin resistance. Our results point to a combination of two major mechanisms to effect nisin resistance: elevating the net charge of the cell wall concomitantly with increasing the thickness of the septum, where the majority of the lipid II molecules reside. Both strategies effectively hamper binding of nisin to lipid II molecules in the bacterial cytoplasmic membrane, preventing abduction of lipid II from sites of cell-wall synthesis and preventing cell lysis by pore formation.

**METHODS**

**Bacterial strains, growth conditions and MIC determinations.** The strains used in this study were L. lactis IL1403 (Chopin et al., 1984), L. lactis IL1403 NisR (Kramer et al., 2004) and L. lactis MG1363 ΔdltD (Dwat et al., 1997). All strains were grown at 30 °C in M17 (Terzaghi & Sandine, 1975) broth with 0.5 % glucose (GM17) and antibiotics where appropriate (chloramphenicol, 5 μg ml⁻¹; nisin, 3000 μg l⁻¹; when using the NisR strain of L. lactis). GM17 plates contained 1.5 % agar. The MIC was determined as described previously (Kramer et al., 2004).

**Isolation of a NisR variant of L. lactis IL1403.** A nisin stock solution was derived from nisinaplin (2.5 % nisin; Aplin and Barrett) as described by Pol & Smid (1999). A NisR isogenic variant of L. lactis IL1403 was obtained as described previously (Kramer et al., 2004). This strain was used for all experiments.

**LTA purification and structure analysis.** LTA purification was performed by culturing 5 l each of L. lactis IL1403, L. lactis IL1403 NisR and L. lactis MG1363 ΔdltD overnight at 30 °C in 500 ml flasks in GM17. Twenty grams (wet wt) of cells from each strain were harvested, resuspended in 60 ml 0.1 M sodium citrate buffer, pH 4.8, and disrupted with a Branson sonifier (250 W) for 5 min at an output level of 6. Between every cycle the cells were kept on ice for 5 min. LTA purification was done essentially as described previously (Morath et al., 2001). Briefly, the disrupted cells were mixed with 60 ml n-butanol (Merck), stirred for 30 min at room temperature and centrifuged at 13 000 g for 40 min at room temperature. The centrifugation step was lengthened to obtain a higher yield of LTA. The aquatic phase was lyophilized, filtered using a 0.22 μm NC 20/20 membrane filter (Schleicher and Schuell), resuspended in 5 ml chromatography start buffer (15 % n-propanol in 0.1 M ammonium acetate, pH 4.7) and centrifuged at 45 000 g for 15 min (Sorvall rotor, SS34) at room temperature. The supernatant was subjected to FPLC on octyl-Sepharose CL 4B (Amersham Pharmacia Biotech) with a 15–60 % propanol gradient. The phosphat content of LTA was estimated according to the protocol of Rouser
Electron microscopy. Overnight cultures of either *L. lactis* IL1403 or *L. lactis* IL1403 Nis<sup>R</sup> were harvested at room temperature by centrifugation for 1 min at 10000 g. Cells were fixed in 3 % glutaraldehyde in 0.1 M sodium cacodylate buffer, pH 7.2 (CAC), and kept on ice for 2 h. Subsequently, the cells were washed three times in CAC, fixed in 1 % osmium tetroxide and 5 % potassium bichromate (1/1) in CAC at room temperature, and washed three times with MilliQ water. Contrast colouring was performed with 0.4 % uranyleacetate, and rehydration with a 100, 96, 70 and 50 % ethanol solution series. The cells were embedded in Epon synthetic resin (Resolution Europe) and sliced on a cryo ultramicrotome (Reichert-Jung). The sliced cells were examined using a Philips CM10 electron microscope (FEI Company).

Subcellular localization of nisin and vancomycin at the *L. lactis* surface. Carboxyfluorescent-labelled nisin and vancomycin were isolated essentially as described previously (Hasper *et al.*, 2006). A culture of *L. lactis* grown overnight was diluted 1 : 30 and grown under the appropriate growth conditions for about 3 h to an OD<sub>600</sub> of about 0.8. Subsequently, fluorescein-labelled vancomycin (1000 μg l<sup>−1</sup>) or fluorescein-labelled nisin (200 μg l<sup>−1</sup>) was added to 0.5 ml culture and the cells were incubated under the appropriate growth conditions for another 15 min. The culture was centrifuged, then the cell pellet was washed in PBS (10 mM potassium phosphate, pH 7.5, 150 mM NaCl), fixed in 1.6 % formaldehyde in PBS and kept on ice for 1 h. After three washes in PBS, the cells were resuspended in PBS and an equal volume of 1 % low-melting-point agar was added for fixation. Cells were inspected with a Zeiss confocal laser scanner (CLSM) 510 (excitation wavelength 488 nm, detection between 500 and 550 nm).

RESULTS

Susceptibility testing of a Nis<sup>R</sup> variant of *L. lactis*

A Nis<sup>R</sup> variant of *L. lactis* IL1403 was used that was able to grow on media containing 3000 μg nisin l<sup>−1</sup>, which is 75 times the MIC of the parental strain (40 μg l<sup>−1</sup>; Kramer *et al.*, 2004). Vancomycin is another antimicrobial substance that inhibits cell-wall synthesis by binding to lipid II. This glycopeptide antibiotic binds to lipid II at the L-Lys-D-Ala-D-Ala terminus of the pentapeptide side chain (Sheldrick *et al.*, 1978), but it does not make pores. Vancomycin can be used as an indicator of the location of lipid II in the cytoplasmic membrane of nisin-sensitive and nisin-resistant strains (Daniel & Errington, 2003). However, as these authors also show, a significant proportion of the vancomycin (~50 %) was bound to free D-Ala-D-Ala termini in the cell wall. It has been shown that the peptidoglycan layer of *L. lactis* contains a significant proportion of free D-Ala-D-Ala termini (25 % of intact pentapeptides; Courtin *et al.*, 2006). Thus, vancomycin binding to free D-Ala-D-Ala termini will contribute significantly to the signal. Sensitivity to vancomycin was tested in the Nis<sup>R</sup> strain as well as in strain *L. lactis* MG1363 ΔDltD which is five times more sensitive to nisin than the wild-type strain, probably due to changes in D-alanylation of LTA. No cross-resistance between nisin and vancomycin was observed in *L. lactis* IL1403 Nis<sup>R</sup> and *L. lactis* MG1363 ΔDltD. The MIC value for vancomycin was unchanged for all the strains investigated. Vancomycin resistance has been shown to be caused by a decrease in the degree of cross-linking (resulting in an increase in free D-Ala-D-Ala termini) together with a thicker cell wall (Cui *et al.*, 2000). This implies that the total amount of free D-Ala-D-Ala termini present in the cell wall does not change significantly with the level of resistance to nisin.

Vancomycin and nisin bind at or close to cell-division sites in *L. lactis*

The total lipid II content in the Gram-positive bacterial cytoplasmic membrane is not correlated with nisin sensitivity (Kramer *et al.*, 2004). Possible changes in the subcellular location of lipid II in the cytoplasmic membrane of *L. lactis* wild-type and Nis<sup>R</sup> strains were investigated, using direct binding of carboxyfluorescein-labelled vancomycin (Daniel & Errington, 2003). This was done to determine whether the binding pattern of nisin to lipid II is altered in the Nis<sup>R</sup> strain. As described above, vancomycin not only binds to the lipid II precursor, but also binds to D-Ala-D-Ala termini present in the peptidoglycan. Thus, as well as localizing lipid II and native peptidoglycan (Daniel & Errington, 2003), carboxyfluorescein-labelled vancomycin will stain other parts of the cell wall. Approximately equal amounts of vancomycin were bound to wild-type *L. lactis*, *L. lactis* Nis<sup>R</sup> and *L. lactis* ΔDltD (Fig. 1a), indicating that the lipid II content in the cytoplasmic membrane is similar in all three strains. As described for *S. pneumoniae* (Daniel & Errington, 2003), vancomycin was predominantly present at the cell-division sites in all three strains of *L. lactis* (Fig. 1a). Approximately equal amounts of vancomycin were bound at newly formed cell-division sites in *L. lactis* ΔDltD, compared to wild-type *L. lactis* and *L. lactis* Nis<sup>R</sup> cells (Fig. 1a).

A quite different pattern was observed for binding of carboxyfluorescein-labelled nisin to lipid II in the cytoplasmic membranes of *L. lactis* wild-type, Nis<sup>R</sup> and ΔDltD cells. In this case, asymmetric fluorescent patches were visible in the bacterial membrane (Fig. 1b). In sensitive cells, the nisin–lipid II patches were located near possible (new) cell-division sites. These patches were also observed in the rod-shaped *Bacillus subtilis*, where lipid II is abducted from its original location to form nisin–lipid II patches (Hasper *et al.*, 2006). Substantially less nisin was bound to cells of *L. lactis* Nis<sup>R</sup> than to cells of its parent, although the binding pattern was basically identical to that observed in the parent strain (Fig. 1b). This raises the question as to which characteristics of the cell wall might cause decreased binding of nisin to the cellular membrane and lipid II. The binding pattern of *L. lactis* ΔDltD was also essentially the same as that of the parent (Fig. 1b).
The cell wall in *L. lactis* NisR is thickened at the cell-division site

Since the nisin–lipid II patches appear to be formed near the (future) cell-division sites (see Fig. 1a), this suggests that *L. lactis* is most sensitive to nisin at this location. In the NisR strain, the acquisition of nisin resistance could be associated with molecular and/or structural changes in the cell wall. Since earlier studies have reported that nisin resistance in *Listeria innocua* is accompanied by an increased overall thickness of the peptidoglycan (Maisnier-Patin & Richard, 1996), the cell walls of *L. lactis* and the NisR derivative were examined by electron microscopy (Fig. 2a, b). The overall cell-wall thickness was 17 ± 2 nm, as measured in 10 different cells, in both the NisR and the parental strains. Notably, a dramatic difference in thickness of the peptidoglycan was observed specifically at the cell-division sites: the cell-division site of *L. lactis* NisR (11 ± 2 nm) was at least twice as thick as that of *L. lactis* (5 ± 2 nm). Thus, upon the acquisition of nisin resistance, the cell wall of *L. lactis* NisR thickens at the sites where lipid II is primarily located in the cell.

LTA is less negatively charged in *L. lactis* NisR

The fact that *L. lactis* ΔdltD is more sensitive to nisin than its parent suggests that LTAs play an important role in the acquired nisin resistance mechanism. LTA was isolated from *L. lactis*, *L. lactis* NisR and *L. lactis* ΔdltD, purified by FPLC and analysed by NMR spectroscopy. The NMR spectra show the resonance signal intensities of the D-alanine, galactose, glycerol and fatty acid fractions of the purified LTAs, which allows quantification of the substituents and to determine the mean glycerophosphate chain length (Morath et al., 2001). LTA of *L. lactis* NisR contains more D-alanyl esters and more galactose than the parent strain (Table 1), and the backbone length is shortened in *L. lactis* NisR. LTA of *L. lactis* ΔdltD contains almost no D-alanine, but exhibits a fourfold increase in galactose substitution relative to the wild-type strain. In conclusion, LTA of *L. lactis* NisR is shorter in length and contains more D-alanine and galactose than its sensitive counterparts, making the cell-wall barrier less negatively charged.

### DISCUSSION

Lipid II is the docking molecule for the lantibiotic nisin (Breukink et al., 1999). We have shown previously that the lipid II content in the bacterial cytoplasmic membrane is not directly correlated with nisin sensitivity of the bacterium (Kramer et al., 2004). At least three mechanisms for nisin resistance in bacteria can be envisaged. First, an increase in the turnover of the sugar moieties to form peptidoglycan may take place, which would decrease the possibility of nisin binding to lipid II. Second, other molecules in the cytoplasmic membrane may prevent nisin from binding to the lipid II molecule. Third, nisin could be prevented from reaching the cytoplasmic membrane, and thus lipid II, due to changes in the cell wall such as an increase in thickness and/or a change in overall charge. A difference in charge would not affect sensitivity of the bacterium to vancomycin, since vancomycin is uncharged.

Earlier studies indicated that genes involved in cell-wall synthesis were involved in the acquisition of nisin resistance (Kramer et al., 2006). Furthermore, it is possible that changes in the subcellular location of the lipid II molecule make cells less susceptible to nisin. The location...
of lipid II in the cytoplasmic membrane was visualized using fluorescently labelled vancomycin, an antibiotic that binds to D-Ala-D-Ala termini present in peptidoglycan, and precursors thereof (such as lipid II). Fluorescently labelled vancomycin was mainly localized at the septum in \( L. \textit{lactis} \), \( L. \textit{lactis} \text{Nis}^R \) and \( L. \textit{lactis} \text{Adh}^D \). This result shows that the cell-wall synthesis machinery, including lipid II, is located at the cell-division sites in \( L. \textit{lactis} \), as also observed in the coccoid \( S. \textit{pneumoniae} \), as well as in the rod-shaped bacteria \( B. \textit{subtilis} \) (Daniel & Errington, 2003; Hasper \textit{et al.}, 2006) and \( B. \textit{megaterium} \) (Hasper \textit{et al.}, 2006).

Binding studies with fluorescently labelled nisin show that \( L. \textit{lactis} \text{Nis}^R \) binds significantly less nisin than \( L. \textit{lactis} \), whereas the binding pattern seems to be the same. In contrast to vancomycin binding, nisin and lipid II appear to form large complexes, which are visible as patches close to cell-division sites in the fluorescent images, rather than being distributed over the entire cell-division site. Abduction of lipid II and formation of giant complexes was also observed with fluorescently labelled nisin in \( B. \textit{subtilis} \) and \( B. \textit{megaterium} \) and with giant unilamellar vesicles containing lipid II (Hasper \textit{et al.}, 2006).

To determine how \( L. \textit{lactis} \text{Nis}^R \) is able to shield the cytoplasmic membrane and lipid II from nisin attack, the cell wall was examined in \( L. \textit{lactis} \) wild-type and its \( \text{Nis}^R \) derivative by electron microscopy. A locally thickened cell wall at the septum was observed. In \( L. \textit{lactis} \text{Nis}^R \), this is the primary location of lipid II in the cell. Thickening of the peptidoglycan was also observed at possible future cell-division sites. Localized thickening of the cell wall can be

---

**Fig. 2.** Electron microscopic analysis of \( L. \textit{lactis} \) (a) and \( L. \textit{lactis} \text{Nis}^R \) (b). Representative cells are shown. The overall cell-wall thickness is 17 nm in both strains. The thickened septa in the \( \text{Nis}^R \) strains are clearly visible at different stages of division in the enlarged images on the left (a) and the right (b).
the result of an increased turnover of the lipid II headgroup. This would decrease the lifetime of lipid II and reduce the amount of binding of nisin to lipid II. An increased turnover of sugar moieties in the lipid II molecule could be due to a higher expression of certain penicillin-binding proteins (PBPs), which are involved in peptidoglycan chain cross-linking and strand elongation (van Heijenoort, 2001). Indeed, in L. lactis NisR, expression of a gene encoding PBP2A is increased relative to L. lactis (Kramer et al., 2006). A putative PBP has also been shown to be more highly expressed in a NisR strain of Listeria monocytogenes (Gravesen et al., 2001). It has also been observed that PBP1A plays an essential role in resistance to cationic antimicrobial peptides in group B Streptococcus (Hamilton et al., 2006). However, it was not established whether the cell wall was (locally) thickened. Alternatively, nisin could be less effective in abducting lipid II molecules to form patches in the NisR strain because of the lower accessibility to lipid II due to the thickened peptidoglycan at the septa. The observed localized thickening of the cell wall at only the septa in L. lactis NisR is in contrast with observations in Listeria innocua, where the entire cell wall is thicker in a NisR strain (Maisnier-Patin & Richard, 1996). Listeria innocua is a fairly nisin-insensitive bacterium and it is therefore difficult to use this species in a comparison of resistance development in L. lactis, since L. lactis is very sensitive to nisin. Thickening of the entire cell wall could be a last resort for Listeria innocua to protect itself from the action of nisin. Recently, it was reported that immediate lethal doses of nisin caused membrane permeabilization, followed by accelerated cell division, cell-envelope biosynthesis inhibition and aberrant cell morphogenesis in B. subtilis (Hyde et al., 2006). This is in line with our findings that resistant strains appear to behave in the opposite way: the cell division time for the NisR strain is longer than in the wild-type strain, which indicates slower cell division (data not shown), thereby making lipid II less accessible.

LTA is the major determinant of cell-wall electrostatic interactions, because it is predominantly negatively charged. In addition, it can contain positively charged d-alanyl esters. A large difference was observed between the LTA of L. lactis and that of L. lactis NisR: the latter contains many more d-alanyl esters and more galactose. This is in accordance with transcriptome analysis data, showing higher expression of genes of the dlt and gal operons (Kramer et al., 2006), and is also in accordance with data from a strain of Staphylococcus aureus containing several copies of the dlt operon. The dlt operon is responsible for d-alanyl ester substitution in LTA, leading to the incorporation of positive charges. The Staphylococcus aureus strain with increased dlt operon expression was less sensitive to various antimicrobial peptides, including nisin (Peschel et al., 1999). Nisin has a net positive charge and the increase of positive charge in the cell wall probably hampers nisin from reaching lipid II in the cytoplasmic membrane by electrostatic repulsion. In a Lactobacillus plantarum strain, LTA showed higher levels of glucose when the dlt operon was deleted (Palumbo et al., 2006). This suggests that by deleting the dlt operon, other sugar moieties increase in number in the LTA. It has also been described in several studies that deletion of dltA in Enterococcus faecalis and Streptococcus pneumoniae, resulting in the absence of D-alanine in the teichoic acids, produces a strain that is more sensitive to antimicrobial peptides (Kovacs et al., 2006; Fabretti et al., 2006). However, these strains are less sensitive to nisin than L. lactis. It has been suggested that many low-G+C Gram-positive bacteria have teichoic acids that contain D-alanine residues in order to protect themselves from the action of antimicrobials such as nisin (Kovacs et al., 2006). An increase in substituted LTA might make the cell wall more tightly packed, thereby shielding the membrane and lipid II from nisin binding. It is also interesting to note that the backbone length of LTA is significantly reduced in L. lactis NisR, which could affect cell-wall packing density. In contrast to LTA of NisR Streptococcus bovis, L. lactis NisR cells did not contain more LTA (Mantovani & Russell, 2001), since the same amount of LTA was purified from wild-type cells as from L. lactis NisR cells. The observed increased sensitivity of L. lactis dltD can therefore be attributed to an increase in negative charge in the LTA, which might lead to a more efficient passage of nisin through the cell wall to reach lipid II and form pores. The acquired NisR phenotype observed in L. lactis is most probably a combination of at least two main effects that add up to a final high nisin resistance level. Shielding of lipid II by the cell wall is acquired by changing the overall charge of LTA. The reduction in accessibility is also accomplished by local thickening of the cell wall at the cell-division sites, where lipid II is most abundant. Shielding of the membrane, and thus the lipid II molecule, appears to

<table>
<thead>
<tr>
<th>Strain</th>
<th>Galactose in LTA (%)</th>
<th>D-Alanine in LTA (%)</th>
<th>Non-substituted LTA (%)</th>
<th>Backbone length</th>
</tr>
</thead>
<tbody>
<tr>
<td>L. lactis II1403</td>
<td>11.8 ± 1.6</td>
<td>28.5 ± 6.6</td>
<td>59.8 ± 5.2</td>
<td>16</td>
</tr>
<tr>
<td>L. lactis II1403 NisR</td>
<td>25.0 ± 5.2</td>
<td>45.0 ± 6.0</td>
<td>30.0 ± 1.8</td>
<td>7</td>
</tr>
<tr>
<td>L. lactis MG1363 ΔdltD</td>
<td>46.0 ± 0.4</td>
<td>5.9 ± 3.5</td>
<td>48.0 ± 3.3</td>
<td>12</td>
</tr>
</tbody>
</table>

Table 1. Substitutions in LTA of L. lactis strains

LTA was isolated from L. lactis, L. lactis NisR and L. lactis ΔdltD, purified by FLPC and analysed by NMR spectroscopy, as described in Methods. The NMR spectra (see Fig. S1, available with the online version of this paper) were used to quantify the substituents in relation to the phosphate content and to determine the mean glycerophosphate chain length.
be the major defence mechanism of *L. lactis* IL1403 against nisin.

**ACKNOWLEDGEMENTS**

We gratefully acknowledge Leo Cobianchi and Klaas Sjollema for the excellent technical assistance in LTA isolation and electron microscopy of *L. lactis*. This work was supported by the NWO and STW grant 349–5257. Project title: Lipid II, the target for the lantibiotic nisin.

**REFERENCES**


Edited by: D. A. Mills