N-Deacetylation of *Sinorhizobium meliloti* Nod Factors Increases Their Stability in the *Medicago sativa* Rhizosphere and Decreases Their Biological Activity

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Accepted 24 September 1999.

Nod factors excreted by rhizobia are signal molecules that consist of a chitin oligomer backbone linked with a fatty acid at the nonreducing end. Modifications of the Nod factor structures influence their stability in the rhizosphere and their biological activity. To test the function of *N*-acetyl groups in Nod factors, NodSm-IV(C_{16:2},S) from *Sinorhizobium meliloti* was enzymatically *N*-deacetylated in vitro with purified chitin deacetylase from *Colletotrichum lindemuthianum*. A family of partially and completely deacetylated derivatives was produced and purified. The most abundant chemical structures identified by mass spectrometry were GlcN(C_{16:2})-GlcNAc-GlcNH₂-GlcNAc(OH)(S), GlcN(C_{16:2})-GlcNAc-GlcNH₂-GlcNH₂(OH)(S), and GlcN(C_{16:2})-GlcNH₂-GlcNH₂-GlcNH₂(OH)(S). In contrast to NodSm-IV(C_{16:2},S), the purified *N*-deacetylated derivatives were stable in the rhizosphere of *Medicago sativa*, indicating that the *N*-acetyl groups make the carbohydrate moiety of Nod factors accessible for glycosyl hydrolases of the host plant. The *N*-deacetylated derivatives displayed only a low level of activity in inducing root hair deformation. Furthermore, the *N*-deacetylated molecules were not able to stimulate Nod factor degradation by *M. sativa* roots, a response elicited by active Nod factors. These data show that *N*-acetyl groups of Nod factors are required for biological activity.

Additional keyword: lipo-chitooligosaccharides.

Rhizobia interact with their leguminous host plants, e.g., *Sinorhizobium meliloti* establishes nitrogen-fixing symbiosis with *Medicago sativa* (alfalfa) (Bladergroen and Spaink 1998; Cohn et al. 1998; Schultze and Kondorosi 1998). Legume roots secrete (iso)flavonoids that induce the synthesis of spe-

cific nodulation signals called Nod factors (NFs). They are modified lipo-chitooligosaccharides, i.e., β -1,4-linked oligomers of *N*-acetyl-D-glucosamine, with a fatty acid replacing the *N*-acetyl group on their nonreducing end (Long 1996; Promé 1996). Structural modifications of NFs at the reducing or nonreducing end have been shown to be important determinants of host specificity. For example, the sulfate group on the reducing end of NFs from *S. meliloti* is required for nodulation of *Medicago sativa*, while mutant strains producing non-sulfated NFs gain the ability to form nodules on *Vicia sativa*, a nonhost plant of *S. meliloti* (Roche et al. 1991). Based on the examination of plant responses to NFs, such as root hair deformation, it is generally thought that structural modifications of NFs influence their binding to putative plant receptors. Moreover, these modifications may affect the stability of NFs in the rhizosphere of the host plant and protect their carbohydrate moiety against degradation and inactivation by plant chitinases and other glycosyl hydrolases (Staehelin et al. 1994a, 1994b; Minic et al. 1998; Schultze et al. 1998). In *M. sativa* roots, NFs are hydrolyzed by a specific "lipo-disaccharide forming" NF hydrolase. The activity of this glycosyl hydrolase is stimulated by NFs from *S. meliloti* at nanomolar concentrations, indicating that NFs induce their rapid degradation in the rhizosphere of the host plant (Staehelin et al. 1995, 1997).

The oligosaccharide core structure of rhizobial NFs is synthesized by NodC, an *N*-acetylglucosaminyltransferase, that has sequence homologies to chitin synthases (Geremia et al. 1994; Kamst et al. 1999). This suggests that a core structure of *N*-acetylglucosamine oligomers is essential for the biological activity of NFs. However, the requirement of the *N*-acetyl groups has not been experimentally proven yet. To test the function of the *N*-acetyl groups in NFs, we prepared derivatives from the *S. meliloti* NodSm-IV(C_{16:2},S) that have glucosamine instead of *N*-acetylglucosamine molecules in their carbohydrate core structure. These molecules were obtained by an in vitro *N*-deacetylation reaction with a purified chitin deacetylase from the plant pathogen *Colletotrichum lindemuthianum* (Tokuyasu et al. 1996). Chitin deacetylases are enzymes that deacetylate the polymer chitin and its oligosaccharides (e.g., Trudel and Asselin

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1990; Kafetzopoulos et al. 1993; Alfonso et al. 1995; Tokuyasu et al. 1996, 1997). The N-deacetylated NF derivatives were purified, analyzed by mass spectrometry, and characterized with respect to their stability in the rhizosphere and their biological activity on the host plant *M. sativa*.

Fig. 1. Chemical structures. **A**, NodSm-IV($C_{16:2}$,S), the tetrameric and sulfated Nod factor from *Sinorhizobium meliloti* was used as substrate for the deacetylation reaction. **B**, Structure of the doubly N-acetylated, mono-N-acetylated, and completely deacetylated derivatives identified by MS/MS. R = $C_{16:2}$ acyl chain.

RESULTS

Deacetylation of NodSm-IV($C_{16:2}$,S) and purification of its N-deacetylated derivatives.

To test the requirement of the N-acetyl groups in NFs, we enzymatically prepared N-deacetylated NF derivatives. NodSm-IV($C_{16:2}$,S), a tetrameric, sulfated NF purified from *S. meliloti* (Figs. 1A and 2A) was incubated with purified chitin deacetylase from the fungus *C. lindemuthianum*. This enzyme has the capacity to release N-acetyl groups from chitin and chitin oligomers as well (Tokuyasu et al. 1997). As shown in Figure 2, NodSm-IV($C_{16:2}$,S) was a substrate for the chitin deacetylase. After incubation, the reaction mixture was analyzed by C_{18} reverse-phase high-pressure liquid chromatography (HPLC). Incubation of NodSm-IV($C_{16:2}$,S) for 1 h resulted in several overlapping new peaks that showed a delayed retention time on the HPLC chromatograms (Fig. 2B). After 3 h, solely N-deacetylated derivatives were seen, while the substrate NodSm-IV($C_{16:2}$,S) was not detected anymore (Fig. 2C).

Figure 3 summarizes the purification procedure that was used to separate the N-deacetylated derivatives from traces of remaining, intact NodSm-IV($C_{16:2}$,S). After incubation of the NF with chitin deacetylase, the reaction mixture was acidified, loaded on a Dowex column, and finally eluted with 2% NH_3 . A separate experiment confirmed that the intact NodSm-IV($C_{16:2}$,S) was not able to bind to the column under these conditions and that a mixture of NodSm-IV($C_{16:2}$,S) and its N-deacetylated derivatives could be separated by this method (data not shown). After this purification step, the N-deacetylated reaction products were purified by reverse-phase HPLC (35% acetonitrile/water, 40 mM ammonium acetate). Finally, the preparation was desalted with a Polygosil column and called purified "material 1."

Material 1 did not contain detectable amounts of intact NodSm-IV($C_{16:2}$,S) as analyzed by HPLC chromatograms and mass spectrometry (data not shown). However, it was not excluded that the purified preparation still contained NodSm-IV($C_{16:2}$,S) in very low amounts. Therefore, an additional purification step to eliminate NodSm-IV($C_{16:2}$,S) from the N-deacetylated molecules was achieved with intact roots of the host plant *M. sativa*. We had shown previously that young *M. sativa* roots were able to rapidly hydrolyze NFs of *S. meliloti* (Staehelin et al. 1994b) and that NodSm-IV($C_{16:2}$,S) was degraded in the rhizosphere by an extracellular "lipodisaccharide forming" NF hydrolase (Staehelin et al. 1995; see also Figure 4). Moreover, we observed that active NFs were strongly bound to the root when nanomolar concentrations of NFs were added to the rhizosphere (C. Staehelin and M. Schultze, unpublished results). Hence, roots of the host plant *M. sativa* have a strong NF degradation activity and binding affinity for NodSm-IV($C_{16:2}$,S). Taking advantage of this, incubation of the N-deacetylated NF derivatives (material 1) with intact *M. sativa* roots was applied as a method to eliminate the possible trace amounts of NodSm-IV($C_{16:2}$,S). When the N-deacetylated NF derivatives were incubated with intact roots for 24 h, only small amounts of a hydrolytic degradation product were observed on HPLC chromatograms (Fig. 2E). This molecule, released from partially N-acetylated molecules, was identified by mass spectrometry as acetylated disaccharide GlcN($C_{16:2}$)-GlcNAc(OH) with a pseudo-molecular ion at $m/z = 617$ and a fragment ion of $m/z = 396$

(data not shown). Most of the material, however, was not degraded by the *M. sativa* root. These nonhydrolyzed molecules present in the incubation medium were purified from the lipo-disaccharide by reverse-phase HPLC, desalted, and called purified "material 2" (Figs. 2D and 3).

Analysis by mass spectrometry.

All fractions were analyzed by liquid secondary ion mass spectrometry (LSIMS). In the negative ionization mode, the presence of the deprotonated molecular ions $(M-H)^-$ of each species was expected. Both preparations (material 1 and material 2) exhibited mostly two peaks at m/z 1059.5 and 1017.6 (Fig. 5). The first one corresponded to mono-N-deacetylated NodSm-IV($C_{16:2},S$) (molecular weight 1060.5), and the second peak was due to doubly N-deacetylated NodSm-IV($C_{16:2},S$) (molecular weight 1018.6). In the positive ion mode, protonated molecules were generated, together with several fragment ions. The same molecules as above were identified with MH^+ ions at m/z 1061.4 and 1019.4, respectively, but a third one appeared with MH^+ at m/z 977.4 (Fig. 5). It was attributed to completely N-deacetylated NodSm-IV($C_{16:2},S$) (molecular weight 976.4). All protonated molecules were accompanied by ions shifted down by 80u (loss of SO_3) and by several fragments of the oligosaccharide backbone (Figs. 5 and 6). The molecular ion ratios of these different molecules were very different depending on the ionization mode. This was not surprising, as the different molecular species differed in their net electric charge. Thus, it was not possible to estimate their relative abundance from the mass spectra of the mixture.

To locate the position of the free amino groups in these partially N-deacetylated molecules, MS/MS (tandem mass spectrometry) spectra of the corresponding MH^+ ions were recorded. Main fragmentations occurred at the glycosidic bonds giving B ion series. The MS/MS spectrum of the MH^+ ion at m/z 977.4 (not shown) was consistent with the expected structure of completely N-deacetylated NodSm-IV($C_{16:2},S$). Indeed, the main fragment intervals from the reducing end were 80u (loss of SO_3), 179u (loss of glucosamine), two times 161u (loss of anhydroglucosamine). This series ended at m/z 396, which is the B_1 ion at the nonreducing end bearing the $C_{16:2}$ N-acyl substituent.

Removal of two N-acetyl groups may give rise to three different products. The MS/MS spectrum of the corresponding MH^+ ion at m/z 1019.4 (Fig. 6) showed (from the reducing end) a 80u interval (SO_3), then either 179u ($GlcNH_2OH$) or 221u ($GlcNAcOH$), the former being prominent. The upper

Fig. 2. Analysis of NodSm-IV($C_{16:2},S$) and its N-deacetylated derivatives by reverse-phase high-pressure liquid chromatography. **A**, NodSm-IV($C_{16:2},S$), the Nod factor purified from *Sinorhizobium meliloti*. **B**, Formation of more slowly migrating, N-deacetylated derivatives after incubation of NodSm-IV($C_{16:2},S$) with chitin deacetylase for 1 h. **C**, Same preparation incubated for 3 h. **D**, Purified, N-deacetylated derivatives (material 2). **E**, Formation of the lipo-disaccharide $GlcN(C_{16:2})-GlcNAc(OH)$ from N-deacetylated derivatives (material 1) after incubation with intact *Medicago sativa* roots for 24 h. **F**, Purified N-deacetylated derivatives (material 2) after incubation with intact *M. sativa* roots for 42 h. Degradation products were not detectable. Peak 1: NodSm-IV($C_{16:2},S$), purified from *S. meliloti*. Peak 2: N-deacetylated derivatives from NodSm-IV($C_{16:2},S$). Peak 3: $GlcN(C_{16:2})-GlcNAc(OH)$ separated into its anomers.

intervals from the B_1 ion at m/z 396 were 203u, mostly (anhydro GlcNAc) or 161u (anhydro GlcNH₂). This interpretation indicated that the remaining *N*-acetyl group was partly on the reducing end and partly on the third residue from the reducing end, the most intense ions corresponding to the latter.

Fig. 3. Purification of *N*-deacetylated Nod factor derivatives called "material 1" and "material 2," respectively. Material 2 was obtained after incubation of material 1 with intact *Medicago sativa* roots, followed by a re-purification procedure.

Fig. 4. Stability of NodSm-IV(C_{16:2},S) and its *N*-deacetylated derivatives in the rhizosphere of *Medicago sativa*. NodSm-IV(C_{16:2},S) or its *N*-deacetylated derivatives (material 1) were incubated with intact *M. sativa* roots for different times. After incubation, samples were fractionated on reverse-phase high-pressure liquid chromatography and the ratio of substrate to the lipo-disaccharide GlcN(C_{16:2})-GlcNAc(OH) product was determined. Each point represents the average value obtained from three seedlings.

From this spectrum, it could not be decided whether a *N*-acetyl group was also present on the second residue or not.

The MS/MS spectrum from the doubly *N*-acetylated components (MH⁺ ion at m/z 1061.4) exhibited, between the fragment MH-80 and the subsequent ones, a 221u interval corresponding to *N*-acetylglucosamine, thus locating this residue at the reducing end (Fig. 6). However, a weak ion with a 179u interval indicated that a less abundant component with glucosamine at the reducing end was also present. At the other side, a 203u interval was found between the B_1 ion at m/z 396 and the closer B_2 . A 161u interval, if any, corresponded to a signal with a very weak abundance. Thus, this glucosamine was mostly *N*-acetylated.

Figure 1B shows the chemical structures identified by MS/MS analysis. The most abundant products were the mono-*N*-deacetylated GlcN(C_{16:2})-GlcNAc-GlcNH₂-GlcNAc(OH) (S), the doubly *N*-deacetylated GlcN(C_{16:2})-GlcNAc-GlcNH₂-GlcNH₂(OH)(S), and the completely *N*-deacetylated GlcN(C_{16:2})-GlcNH₂-GlcNH₂-GlcNH₂(OH)(S). Furthermore, two minor components were found: The mono-*N*-deacetylated GlcN(C_{16:2})-GlcNAc-GlcNAc-GlcNH₂(OH)(S) and the doubly *N*-deacetylated GlcN(C_{16:2})-GlcNH₂-GlcNH₂-GlcNAc(OH) (S). Since the polarity and the hydrophobicity of the *N*-deacetylated molecules are very different, it was not possible to exactly determine their relative abundance from the MS/MS spectra.

High stability of *N*-deacetylated NF derivatives in the *M. sativa* rhizosphere.

When incubated with intact *M. sativa* roots, NodSm-IV(C_{16:2},S) was rapidly hydrolyzed and inactivated by a specific "lipo-disaccharide forming" NF hydrolase. It was hypothesized that inactivation of NFs after their perception is an important step of symbiotic signaling (Staehelin et al. 1994b, 1995). Therefore, we tested the stability of *N*-deacetylated derivatives in the *M. sativa* rhizosphere, compared with NodSm-IV(C_{16:2},S). A time course experiment shown in Figure 4 illustrates the difference in stability of *N*-deacetylated derivatives (material 1) compared with intact NodSm-IV(C_{16:2},S) in the rhizosphere of *M. sativa*. Roots of young seedlings were incubated in a solution containing NodSm-IV(C_{16:2},S) or its derivatives for different time periods. After extraction of the medium with *n*-butanol, the substrate and the

Fig. 5. Liquid secondary ion mass spectrometry (LSIMS) spectra of the partially *N*-deacetylated Nod factor derivatives. Molecular region of the positive (top) and negative ion (bottom) LSIMS spectra.

acylated cleavage products were fractionated by reverse-phase HPLC. The graph indicates the percentage of nonhydrolyzed substrate after a given incubation time. The NodSm-IV ($C_{16:2},S$) molecules were rapidly degraded to the lipo-disaccharide GlcN($C_{16:2}$)-GlcNAc(OH), likely by the "lipo-disaccharide forming" NF hydrolase as reported previously (Staehelin et al. 1995). In contrast, only small amounts of GlcN($C_{16:2}$)-GlcNAc(OH) were released from the deacetylated derivatives (material 1) (Fig. 2E). After about 16 h, a plateau was reached and no further hydrolysis was observed (Fig. 4). These findings were confirmed by an additional experiment. After re-purification of the nonhydrolyzed molecules, the material (i.e., material 2) was incubated for a second time with intact roots of freshly prepared *M. sativa* plants. As expected, these molecules were resistant against hydrolysis, even after an incubation time as long as 42 h (Fig. 2F). Similar results were obtained when a mixture of NodSm-IV($C_{16:2},S$) and its N-deacetylated derivatives was tested. NodSm-IV($C_{16:2},S$) was degraded to the lipo-disaccharide, while the deacetylated derivatives remained intact (not shown). These data indicate that the deacetylated derivatives exhibit a high stability in the *M. sativa* rhizosphere and that the *N*-acetyl groups of NFs make the carbohydrate moiety accessible for hydrolases of the host plant.

Low biological activity of N-deacetylated NF derivatives.

NFs are known to induce a number of specific responses on the host plants. To test whether the deacetylated derivatives show a similar inducing activity on *M. sativa*, two different bioassays were used, i.e., the capacity to stimulate root hair deformation and to enhance "lipo-disaccharide forming" NF hydrolase activity in the rhizosphere (Staehelin et al. 1995).

In a first series of experiments, the activity to stimulate root hair deformation on young *M. sativa* roots was determined. As seen in Table 1, the intact NF was able to induce a response with a threshold at 10^{-11} M. When deacetylated derivatives (material 2) were tested, a threshold at about 10^{-7} M was detected, indicating an approximately 10,000-fold lower activity, compared with NodSm-IV($C_{16:2},S$). Excess amounts of deacetylated derivatives were not able to inhibit the root hair deformation responses induced by intact NodSm-IV($C_{16:2},S$), indicating that the deacetylated molecules cannot act as competitive inhibitors (not shown).

In a second series of experiments, the deacetylated derivatives were tested for their ability to stimulate the "lipo-disaccharide forming" NF hydrolase of *M. sativa* (Staehelin et al. 1995). The activity of this enzyme showed a concentration-dependent stimulation when *M. sativa* roots were pretreated with active NFs. For NodSm-IV($C_{16:2},S$), a threshold concen-

Fig. 6. MS/MS spectra of the partially N-deacetylated Nod factor derivatives. MS/MS spectra of the MH^+ ions from doubly (top) and mono (bottom) N-deacetylated species at m/z 1019 and 1061, respectively. The B fragments resulted from glycosidic cleavages with charge retention on the nonreducing part. The Y fragments are their counterpart with charge retention on the reducing end. The mass range was scanned from MH^+ down to m/z 350.

tration of 10^{-8} M was measured (Staehelin et al. 1995). As seen in Figure 7, a pretreatment of roots with 10^{-6} M NodSm-IV(C_{16,2},S) resulted in a fourfold stimulation of the enzyme activity (column 3). The N-deacetylated derivatives (material 2) were not able to stimulate the hydrolase (column 2), i.e., the “lipo-disaccharide forming” activity of pretreated plants was as low as for the control plants, which were mock-treated with Jensen medium (column 1). Addition of excess amounts of N-deacetylated derivatives (material 2) did not influence the effect induced by active NFs (column 4). These results confirm and extend the data obtained from the root hair deformation assay and show that N-deacetylation of NFs dramatically decreases their biological activity.

DISCUSSION

In this paper, we report the preparation of N-deacetylated derivatives of NodSm-IV(C_{16,2},S) with chitin deacetylase from *C. lindemuthianum*. The enzyme was able to release all three N-acetyl groups of the tetrameric NF and produced a family of doubly N-acetylated, mono-N-acetylated, and completely deacetylated molecules. The rhizobial NodB protein involved in NF synthesis, however, specifically N-deacetylates a single acetyl group at the nonreducing end of chitin oligomers, forming mono-deacetylated precursor molecules that can be acylated by NodA protein (John et al. 1993). Despite this difference in substrate specificity, fungal genes encoding chitin deacetylase show remarkable sequence similarities with *nodB* (Kafetzopoulos et al. 1993).

Compared with intact NodSm-IV(C_{16,2},S), the N-deacetylated derivatives displayed an approximately 10,000-fold reduced biological activity in the root hair deformation assay. After incubation with chitin deacetylase, intact NFs were not detected anymore on HPLC chromatograms (Fig. 2C). The following four purification steps further removed trace amounts of NFs from the N-deacetylated derivatives. We assume that each of these steps resulted in an approximately 100-fold reduction of contaminating NodSm-IV(C_{16,2},S). However, we cannot completely exclude that the weak biological activity of the N-deacetylated derivatives in inducing root hair deformation was due to contaminating NodSm-IV(C_{16,2},S). Hence, it is possible that N-deacetylated derivatives are completely inactive in eliciting root hair deformation on *M. sativa* roots. It is worth noting in this context, however, that also the lipo-disaccharide GlcN(C_{16,2})-GlcNAc(OH) or the lipo-trisaccharide GlcN(C_{16,2})-GlcNAc-GlcNAc(OH) retained root hair deformation activity in the micromolar concentration range (Staehelin et al. 1994b).

Table 1. Root hair deformation assay^a

Treatment	Concentration (M)						
	10^{-5}	10^{-6}	10^{-7}	10^{-8}	10^{-9}	10^{-10}	10^{-11}
NodSm-IV(C _{16,2} ,S)	+	+	+	+	+	+	±
N-deacetylated derivatives	+	+	±	–	–	–	–

^a Capacity of NodSm-IV(C_{16,2},S) and its N-deacetylated derivatives (material 2) to induce root hair deformation on *Medicago sativa* was tested at indicated concentrations. Positive and negative scores were given when root hair deformation was present (+) or absent (–), respectively. At the threshold value (±) only a few plants were responsive.

A treatment of roots with excess amounts of N-deacetylated derivatives did not inhibit the responses of the host plant induced by intact NFs. We suggest that N-deacetylated derivatives do not bind to the NF receptors involved in root hair deformation and stimulation of the “lipo-disaccharide forming” NF hydrolase. Hence, these receptors seem to have a strongly reduced affinity for lipo-oligosaccharides when GlcNAc of the core structure is replaced by GlcNH₂.

We have found previously that tetrameric NFs of *S. meliloti* are rapidly degraded by host plants but are highly stable in the rhizosphere of nonhost plants (Staehelin et al. 1994b, 1997). It is tempting to speculate that NF degradation is a prerequisite for a compatible symbiotic interaction and that host plants have evolved specific enzymes that cleave and inactivate the rhizobial signal molecules (Staehelin et al. 1995). In the present study, the release of N-acetyl groups from NFs resulted in a dramatically increased stability in the *M. sativa* rhizosphere. These results indicate that the N-deacetylated derivatives (material 2) are substrates for neither the “lipo-disaccharide forming” NF hydrolase nor other glycosyl hydrolases in the *M. sativa* rhizosphere.

It has been reported by Bec-Ferté et al. (1996) that certain *S. fredii* strains produce various NFs including a compound with glucose inserted to the chitin oligomer chain. This pentameric NF was found to be degradable by a bacterial chitinase; however, its biological activity on roots was not examined. Based on our results with N-deacetylated NF derivatives, we suggest that this glucose-containing NF has low biological activity on host plants.

Fig. 7. Capacity of NodSm-IV(C_{16,2},S) and its N-deacetylated derivatives to stimulate the “lipo-disaccharide forming” Nod factor (NF) hydrolase of *Medicago sativa*. Roots of young *M. sativa* seedlings were pretreated for 19 h with the following solutions: Jensen medium as a control (column 1); 1 μ M N-deacetylated derivatives (material 2) (column 2); 1 μ M NodSm-IV(C_{16,2},S) (column 3); and a mixture of 10 μ M N-deacetylated derivatives with 1 μ M NodSm-IV(C_{16,2},S) (column 4). After incubation, roots were incubated in 5 μ M NodSm-IV(C_{16,2},S) for 2 h. Release of the lipo-disaccharide GlcN(C_{16,2})-GlcNAc(OH) from the NF substrate was analyzed by reverse-phase high-pressure liquid chromatography. Standard deviation of the mean ($n = 3$) is indicated.

In addition to NodSm-IV(C_{16:2}S), we were able to N-deacetylate various other NFs, including the O-acetylated NodSm-IV(C_{16:2}Ac,S) and the nonsulfated derivative NodSm-IV(C_{16:2}) (data not shown). It is possible that all rhizobial NFs are substrates for the chitin deacetylase from *C. lindemuthianum*. The production of N-deacetylated derivatives opens the possibility to reacetylate them chemically by radioactively labeled acetic anhydride, as described for chitosan that can be converted to chitin (e.g., Molano et al. 1977; Boller 1992). Radioactively labeled NFs are valuable tools to characterize NF-binding proteins, which are candidates for putative NF receptors. For in vitro radiolabeling, NFs were chemically modified, e.g., by catalytic hydrogenation, forming less active NFs (Schultze et al. 1992). Alternatively, specific enzymes were required, such as the sulfate transferase NodH (Schultze et al. 1995; Bourdineaud et al. 1995) and the O-acetyl transferase NodL (Bloemberg et al. 1994). Enzymatic N-deacetylation and reacetylation with acetic anhydride, however, would be a general method for in vitro radiolabeling of any NF, without modifying its structure.

MATERIALS AND METHODS

Nod factors and chitin deacetylase.

Purification of the tetrameric and sulfated NodSm-IV (C_{16:2}S) produced by the *S. meliloti* (formerly named *Rhizobium meliloti*) strain 1021(pEK327) has been described previously (Schultze et al. 1992). Chitin deacetylase from a culture filtrate of the fungus *C. lindemuthianum* (ATCC56676) was purified to electrophoretic homogeneity as reported (Tokuyasu et al. 1996), lyophilized in aliquots, and resuspended in H₂O for the enzymatic reaction with NFs.

Deacetylation reaction.

For a typical in vitro deacetylation reaction, 1 mg of NodSm-IV(C_{16:2}S) was incubated with 2 U of purified chitin deacetylase in 0.5 ml of 20 mM sodium phosphate buffer, pH 7.5, at 37°C for 3 h. The deacetylation reaction was monitored by fractionating small aliquots on reverse-phase HPLC under isocratic conditions with 35% acetonitrile/water, 40 mM ammonium acetate as the mobile phase (Staehelin et al. 1994b). NodSm-IV(C_{16:2}S) and the more slowly migrating N-deacetylated derivatives were detected by their absorption at 220 nm. Quantification of the N-deacetylated derivatives was based on their absorption of the C_{16:2} acyl chain at 220 nm.

Purification of N-deacetylated derivatives (material 1).

Step 1: Ion exchange chromatography (according to Drouillard et al. 1997). After incubation, the sample was acidified with 50 µl of 1 M acetic acid and loaded onto a Dowex 50W-X2 column (150 mg dry weight), which was equilibrated with 10 mM acetic acid. As a control, NodSm-IV(C_{16:2}S) was chromatographed under the same conditions. After the columns were washed with H₂O, aliquots were analyzed by reverse-phase HPLC (see above). While NodSm-IV(C_{16:2}S) was found in the flow-through, the charged N-deacetylated derivatives bound to the column and could be eluted with 3 ml of 2 % (vol/vol) aqueous NH₃.

Step 2: Reverse-phase HPLC. The eluted, N-deacetylated derivatives were concentrated under reduced pressure and the residue (a few microliters) was taken up in 100 µl of 35%

acetonitrile/water, 40 mM ammonium acetate. The sample was purified by reverse-phase HPLC as described (Staehelin et al. 1994b) with 35% acetonitrile/water, 40 mM ammonium acetate as the mobile phase. The peak representing the N-deacetylated compounds was collected and 20-fold concentrated under reduced pressure.

Step 3: Desalting step. A C₁₈ column (Machery Nagel, Düren, Germany, Polygosil C₁₈, 60-4063, particle size 40 to 63 µm) was washed with 100% methanol and equilibrated with H₂O. The N-deacetylated compounds were diluted in 1 ml of H₂O and loaded onto the column (200 mg dry weight). The column was washed with 10 ml of H₂O and the deacetylated derivatives were then eluted with 1 ml of 100% methanol and dried in a Speed-Vac evaporator.

Purification of N-deacetylated derivatives (material 2).

Step 4: Incubation in the *M. sativa* rhizosphere. N-deacetylated derivatives (material 1) were incubated with germinated *M. sativa* (cv. Sital) seedlings in 1-ml sterile plastic syringes as described previously (Staehelin et al. 1994b). Briefly, 100 µg of material 1 was taken up in 10 ml of deposit-free Jensen medium containing 0.5 (vol/vol) dimethyl sulfoxide (DMSO). The solution was filled into the plastic syringes (200 µl per syringe) and then a young seedling was placed on top of each syringe. After incubation at 24°C in the dark for 24 h, the seedlings were removed and the contents of the syringes were pooled and extracted with 3 ml of distilled *n*-butanol. The butanol phase was dried in a Speed-Vac evaporator.

Step 2': Reverse-phase HPLC as described for step 2.

Step 3': Samples were desalted as described for step 3.

Analysis by mass spectrometry.

All spectra were recorded on an AutoSpec instrument (Micromass, Manchester, UK) fitted with an LSIMS source and a cesium ion gun. The accelerating voltage was set to 8 kV, the cesium gun to 30 kV. Typically, samples (1 µg or less) were deposited on the LSIMS tip from a water solution and mixed with the appropriate matrix (1:1 glycerol/metanitrobenzyl alcohol, either acidified with 10% trichloroacetic acid for the positive ion mode or alkalized with diisopropylmethyl amine for negative ions). MS/MS spectra were recorded in the constant B/E scanning mode.

Hydrolysis of NodSm-IV(C_{16:2}S) and its N-deacetylated derivatives by *M. sativa* roots.

The in vivo assay with intact *M. sativa* roots was performed as described previously (Staehelin et al. 1994b, 1995). Briefly, roots of *M. sativa* seedlings were incubated in 1-ml plastic syringes filled with 200 µl of Jensen medium containing 0.5% (vol/vol) DMSO and 10 µM NodSm-IV(C_{16:2}S) or its N-deacetylated derivatives, respectively. After incubation at 24°C in the dark, the seedlings were removed. The culture medium of three seedlings was pooled and extracted with an equal volume of *n*-butanol. Substrates and the acylated degradation product GlcN(C_{16:2})-GlcNAc(OH) were fractionated on reverse-phase HPLC with 35% acetonitrile/water, 40 mM ammonium acetate.

Bioassays.

As a bioassay for NF activity, two specific host plant responses were examined, root hair deformation and stimulation

of the "lipo-disaccharide forming" NF hydrolase of *M. sativa*. The root hair deformation assay was performed as reported previously with 20 seedlings for each concentration range (Staehelin et al. 1994b). The activity of the "lipo-disaccharide forming" NF hydrolase was measured as described (Staehelin et al. 1995). Briefly, *M. sativa* seedlings were incubated for 19 h in syringes filled with a given concentration of NFs or N-deacetylated derivatives, respectively. Plants were then transferred to new syringes containing 5 μ M NodSm-IV(C_{16:2},S) and incubated at 24°C in the dark for 2 h. After extraction with *n*-butanol, the substrate and the formed lipo-disaccharide GlcN(C_{16:2})-GlcNAc(OH) were analyzed by reverse-phase HPLC as described (Staehelin et al. 1994b).

ACKNOWLEDGMENTS

This work was partially supported by the Max Planck Research Award to A. K. We appreciated helpful discussions with Peter Mergaert (ISV, CNRS, Gif-sur-Yvette, France).

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