Nitrogen Inhibits Nodulation and Reversibly Suppresses Nitrogen Fixation in Nodules of *Alnus maritima*

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**Abstract.** Symbiotic associations between *Alnus maritima* (Marsh.) Muhl. ex Nutt. (seaside alder) and actinomycetes in the genus *Frankia* Bruchorst result in root nodules in which atmospheric nitrogen (N) is fixed. The economic and environmental benefits of N fixation have led to interest in inducing root nodules during production of *A. maritima*. Because woody plants produced in nurseries typically are provided N fertilizer, our objectives were to determine how applied N influences nodulation of *A. maritima* and to characterize how short-term changes in root-zone N affect the function of nodules. Potted seedlings were grown in perlite that was inoculated with 30 mL of soil from the root zones of mature plants in their native habitat on the Delmarva Peninsula. Each pot was drenched once daily for 10 weeks with nutrient solution that contained ammonium nitrate at 10 concentrations from 0 to 8 mM. Plants that received no ammonium nitrate formed the most nodules, and nodulation decreased linearly as ammonium nitrate increased from 0.25 to 4 mM. Plants treated with ammonium nitrate at 0.5 to 4 mM formed nearly no nodules, while ammonium nitrate at 0.5 mM resulted in vigorous plants with an average nodule count of 70. In a second experiment, a population of nodulated seaside alders was established by irrigating seedlings in inoculated perlite once daily with 0.5-mM ammonium nitrate for 6 weeks. Plants then were provided ammonium nitrate at 0.5, 2, or 4 mM for 2 weeks. Acetylene-reduction assays showed suppressed nodule activity among plants provided 2- and 4-mM ammonium nitrate. Daily irrigation of those plants with N-free solution subsequently led to a rapid depletion of root-zone N and to a concomitant resurgence of nodule activity. These results demonstrate that N fertilization can be managed to promote nodulation of *A. maritima* and show that decreased nodule activity caused by short-term increases in root-zone N is reversible.

*Alnus maritima* is an attractive shrub or small tree native to three small, disjunct provenances in the United States (Schrader and Graves, 2000). Like other alders, *A. maritima* is an actinorhizal species that forms root nodules in which *Frankia* fix gaseous N (Schrader and Graves, 2000; Stibolt, 1978). Unlike most woody plants that cannot benefit directly from atmospheric N, *A. maritima* with functional root nodules might perform well if planted in N-poor soils. We seek to understand how the symbiosis between *A. maritima* and *Frankia* can be established and optimized during the culture of plants in nurseries.

Commercial production of plants with functional root nodules is important for two reasons. First, growers of nodulated *A. maritima* might apply less N fertilizer, which would reduce production costs and the potential for environmental damage caused by run-off of irrigation water. Second, the soils in which *A. maritima* is planted may not contain compatible *Frankia*; installation of nodulated plants would overcome this possible barrier to N fixation in the landscape. Protocols for producing nodulated *A. maritima* are needed. In addition, data are needed on how N fertilization affects the N fixation of nodulated plants, and comparisons of the growth of plants reliant on fixed N and those supplied N fertilizer would help producers assess the practicality of reducing N fertilization.

Formation of functional nodules on actinorhizal plants requires the presence of compatible *Frankia* in edaphic environments conducive to bacterial infection, nodule development, and the activity of nitrogenase. Although numerous physical and chemical factors may govern the nodulation of *A. maritima* and the function of its nodules, N content of the root zone might be the most important during production of plants in nurseries. Researchers working with other alders have demonstrated that high N concentrations restrict nodule formation and activity (Berry and Torrey, 1985; Gentili and Huss-Danell, 2003; Ingestad, 1980; Kohls and Baker, 1989). Huss-Danell et al. (1982) showed that 20-mM ammonium chloride damages the vesicles of *Frankia* and reduces nitrogenase activity in nodules of *Alnus incana* (L.) Moench, an effect that was reversed after ammonium chloride applications were reduced. Burgess and Peterson (1987) suggested using low-N fertilizers to permit nodule formation and N fixation of *Alnus japonica* (Thunb.) Steud. However, no such data have been generated for *A. maritima*.

Our first objective was to define how various concentrations of ammonium nitrate affect the nodulation of *A. maritima* and the subsequent fixation of N. Our second objective was to determine how short-term increases and decreases in applied root-zone N influence N fixation using a population of plants with copious, active nodules. Ammonium nitrate was the N source used to meet both objectives because it often is used commercially. Also, unlike other common N sources, varying the rate at which ammonium nitrate is applied does not directly alter the concentrations of other ions. The results of these experiments enhance our understanding of symbioses of actinorhizal plants and will be useful to growers who want to ensure that their *A. maritima* possess numerous root nodules in which N is fixed at high rates.

**Materials and Methods**

**N effects on establishment of symbioses.** Cold-stratified seeds (*n* = 675) of *A. maritima* ssp. *maritima* indigenous to 38°36'N lat., 75°30'W long. on the Delmarva Peninsula were germinated in clay pots (top diameter = 7 cm, height = 9.5 cm)
filled with coarse perlite and a 3-cm-deep layer of fine-grade vermiculite directly around the seeds. The pots were kept moist with nitrate-free tap water in a greenhouse in which the air temperature was 18 to 30 °C and natural irradiance was supplemented by use of incandescent lamps that provided \(<100 \mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}\) photosynthetically active radiation (PAR) and 16-h photoperiods. Seedlings were fertilized twice daily with 18-mm N from a blend (1:3, by weight) of water-soluble Peters Excel All-Purpose and Peters Excel Cal-Mag (17N–2.2P–13.3K) (Scotts Sierra Horticultural Products, Marysville, Ohio). Research with other alders and our informal observations of \(A.\) \emph{maritima} indicated 18-mm N would prevent root nodulation.

Eighty one-month-old seedlings of uniform size were transplanted from the clay pots to individual plastic pots (top diameter = 12.5 cm, height = 11.5 cm). The medium was coarse perlite to which 30 mL of soil was added per pot as inoculum to provide compatible \emph{Frankia}. The soil was from the root zones of naturally occurring, mature \(A.\) \emph{maritima} on the Delmarva Peninsula and was placed in the upper 2 cm of perlite. Treatments began on 10 Apr. 2003 when the seedlings had two fully expanded true leaves. Each seedling was regarded as an experimental unit, and a randomized complete-block design was used. Eight seedlings were assigned randomly to each of 10 treatments to provide plants with various amounts of N. A quarter-strength, N-free Hoagland solution (Hoagland and Arnon, 1950), modified to provide half of the prescribed Fe, was supplemented with ammonium nitrate at 0, 0.25, 0.5, 0.75, 1.0, 1.25, 1.5, 2.0, 4.0, or 8.0 mM. The pH of all 10 solutions was 5.5. Each plant was irrigated to contain capacity with 300 mL of its prescribed solution on the day of transplanting and inoculation (day 0) and irrigated once daily thereafter with treatment solution. Air temperature in the greenhouse ranged from 23 to 30 °C, and midday PAR was as high as 550 \(\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}\).

After 10 weeks, N fixation of eight randomly selected plants from three ammonium nitrate treatments was estimated using an acetylene reduction assay (Hardy et al., 1968; Huss-Danell, 1978). Each root system was placed in a 1-L mason jar, and airtight seals were created around the stems protruding through the openings in the lids. We removed 100 mL of the atmosphere from each jar and replaced it with 100 mL of ultra-high-purity acetylene. Linearity of ethylene production for over 30 min and undetectable endogenous ethylene from roots after 30 min without acetylene were confirmed in preliminary work. Therefore, a 1-mL sample was removed with a gas-tight syringe after 30 min, reduced to 0.1 mL, and brought back to 1 mL with air. A Varian 3600 Star CX gas chromatograph (Varian, Palo Alto, Calif.) was used to analyze the sample for ethylene. Reduction of acetylene to ethylene by nitrogenase indirectly represents N-fixation activity. The assays were performed in the greenhouse during the middle of the photoperiod. Immediately afterwards, the total number of nodules per plant and the leaf surface area (LI-3100 area meter; LI-COR, Lincoln, Nebr.) of all plants were determined. We expressed nodule counts on a per-plant basis because there was no treatment effect on root dry weight. Plant height was measured, and nodules, stems, leaves, and roots were weighed separately after they were dried at 67 °C for 7 d. Total N content of dried leaves from plants treated with ammonium nitrate at 0, 0.5, 1, 2, and 8 mm was determined. The dependent variables were regressed over the N concentrations applied in irrigation solutions.

**Root-zone N effects on nodule function.** This experiment was conducted in a greenhouse in which the air temperature ranged from 24 to 32 °C and PAR was as high as 590 \(\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}\). Individually potted seedlings were experimental units in a completely randomized design. Phase one of this experiment began with establishing a population of seedlings by using the methods described for the first experiment. Sixty seedlings were transplanted when they were 6 weeks old by using the same materials as in the first experiment. Beginning on 3 July 2003, the 60 pots were irrigated once daily for 7 weeks with a base solution supplemented with 0.5-mm ammonium nitrate, one of the solutions used in the first experiment. This N concentration was selected because it led to vigorous plants with many root nodules during the first experiment. At the conclusion of phase one, we had 60 robust plants that we presumed had numerous root nodules.

Phase two began by randomly assigning 20 of the plants to each of three treatments, which were application of ammonium nitrate at 0.5, 2, and 4 mm once daily. Four plants from each of the three treatments were chosen randomly and harvested after 2 weeks. Acetylene-reduction assays, nodule counts, and plant size were determined as in the first experiment. The 48 remaining plants, 16 in each of the phase-two treatments, entered phase three that day and were irrigated thereafter once daily with the base nutrient solution with no ammonium nitrate. Four plants from each phase-two treatment were harvested after 1, 2, 4, or 8 weeks. Acetylene-reduction assays, nodule counts, and plant size were determined at each harvest. Perlite from pots of all plants harvested after 1 and 2 weeks of phase three was analyzed for total N content. The dry, ground sample was burned in a combustion tube at 950 °C, and exhaust gases were collected. Gaseous N was measured by thermal conductivity with a Leco CHN 2000 (Leco Corp., St. Joseph, Mich.).

**Data analysis.** Data from both experiments were analyzed by using SAS (version 8E; SAS Inst., Cary, N.C.), and analysis of variance was performed by using the general linear model for least square means. Values of \(P > 0.05\) led to the rejection of hypotheses. For experiment two, effects of ammonium nitrate, time, and their interaction were partitioned. Version 8.0 of SigmaPlot (SPSS, Chicago) was used to perform linear regression analyses over time and concentrations of ammonium nitrate.

**Results.**

**N effects on establishment of symbioses.** Mean nodule count for plants that received no ammonium nitrate exceeded 100 and was \(\approx 30\) more than the mean nodule count for plants provided ammonium nitrate at 0.25 mm (Fig. 1). Nodule count decreased linearly as applied ammonium nitrate increased from 0.25 to 4 mm (Fig. 1). The 4-mm treatment led to a mean nodule count of 0.3 per plant, whereas 8-mm ammonium nitrate prevented nodulation (Fig. 1). Acetylene reduction to ethylene was minimal among the nodule-free root systems in the 8-mm treatment, whereas root systems provided no ammonium nitrate yielded the greatest ethylene production at nearly 0.71 \(\mu\text{mol}\cdot\text{h}^{-1}\) per plant during the 30-min assays (Fig. 1). Regressing acetylene reduction per root system across the three ammonium nitrate treatments for which assays were run resulted in a linear relationship: ethylene = 1.1 – 0.34 (ammonium nitrate); \(r^2 = 0.78\). Treatment effects on mean dry weight of nodules per plant were consistent with nodule counts, and dividing nodule masses by nodule counts revealed that the mean dry mass of an individual nodule, 0.77 mg, was similar among treatments. Mean plant height decreased linearly, and mean total N content of leaves increased linearly, as applied ammonium nitrate increased (Fig. 2). Stems of plants that received no ammonium nitrate appeared less stout and erect than stems of...
Nor was leaf surface area affected (when ammonium nitrate is applied at concentrations ≥ 4 mM (Fig. 1). Activity of nodules that form in low-N root zones is sensitive to changes in N. Decreases in activity caused by short-term increases in N are reversed rapidly as root-zone N declines (Fig. 3). Producers of A. maritima can use these data to manage N applications to ensure nodulation, N fixation, and minimal waste of N fertilizers. Our results also show that growers and landscape managers may reverse the deleterious effects of N fixation in inadvertent increases in root-zone N by leaching N from the adjacent substrate.

The N-induced reduction in nodule formation and activity we observed is consistent with previous research on other species of alder. Kohls and Baker (1989) found that nodulation of Alnus glutinosa (L.) Gaertn. is negatively correlated with root-zone N.
concentration and is completely inhibited when N exceeds 2 mm, possibly because of damage to the vesicles of *Frankia* (Huss-Danell et al., 1982). Although the response of *A. maritima* was similar, regression analysis did not predict that 2- to 4-mm ammonium nitrate completely inhibits nodule formation (Fig. 1). This suggests that nodulation of *A. maritima* may be less sensitive to N than is nodulation of *A. glutinosa*, but direct comparisons of the two species are needed. Our results are attributable directly to N because other nutrient elements were not varied and the pH of all solutions was 5.5. In addition, a preliminary study we conducted showed that daily irrigation to container capacity prevented changes >1 pH unit between irrigations. It was important for us to maintain uniform pH among treatments because numerous researchers, beginning with Quispel (1958), have characterized pH effects on nodulation. Berry and Torrey (1985) considered pH 5.5 optimal for nodulation and reported that pH < 4.5 inhibits the formation of nodules on roots of *A. glutinosa*, *Alnus incana* ssp. rugosa (L.) Furlow, and *Alnus rubra* Bong.

Most of the many nodules observed after phase two of our second experiment likely formed as plants received 0.5-mm ammonium nitrate during the first phase, thereby substantiating the effect of 0.5-mm ammonium nitrate documented during the first experiment. In addition, data collected at the end of phase two of the second experiment provide evidence that the activity of nodules formed under low-N conditions is suppressed rapidly upon exposure to increased N. Phase three was designed to test whether this suppression was reversible, and treatment differences no longer existed after daily application of N-free solution for 1 week (Fig. 3). Analysis of the perlite confirmed a rapid leaching of N concomitant with increased nodule activity. Huss-Danell et al. (1982) similarly documented recovery of nodule activity among plants of *A. incana* after a short-term exposure to 20-mm ammonium chloride. Considered collectively, the three phases of the second experiment illustrate the dynamic influence of N during establishment and function of N-fixing symbioses on roots of *A. maritima*. Producers of this species, and managers of *A. maritima* installed in the landscape, should recognize that low N promotes nodulation and that increases in N due to fertilization or other causes will reduce nodule function until N concentrations in the root zone are reduced.

We have reached several important conclusions regarding how N affects symbioses between *A. maritima* and *Frankia*, but several issues remain unresolved. Whether ammonium and nitrate influence the symbioses to the same extent and via similar mechanisms is unclear. How long nodulated plants can be exposed to N concentrations that inhibit N fixation without causing irreversible damage to the symbioses is another relevant issue. Whether it is necessary to inoculate *A. maritima* with *Frankia*, and if so, whether it is critical to provide bacteria from the soils where the species is native are unknown. We occasionally find nodules on uninoculated plants cultured in medium without soil, but data are needed on the ubiquity and functional variability of *Frankia* compatible with *A. maritima*. The conclusions we have drawn regarding N effects can serve as a basis for developing new hypotheses regarding these unresolved issues. Moreover, our findings demonstrate that the dry mass and leaf surface area of *A. maritima* produced with little or no applied N can be similar to those traits of *A. maritima* provided luxurious concentrations of ammonium nitrate if compatible *Frankia* are present in root zones. Although application of no N evoked the most nodules (Fig. 1), we observed that shoot systems of plants in that treatment were shaped irregularly and had leaves that were not as intensely green as the foliage of plants in the other treatments. Therefore, we recommend use of ammonium nitrate at 0.5 to 2 mm to enhance leaf N content and reduce height (Fig. 2) of plants with a large number of active root nodules (Fig. 1).

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