

Effect of nitrate and ammonium on growth of transplanted Norway spruce seedlings: a greenhouse study

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Effects of nitrate and ammonium nitrogen (N) on the growth of two-year-old Norway spruce seedlings (*Picea abies*) were studied after transplanting in a greenhouse. In a preliminary experiment (Exp. 1), seedlings were transplanted into sandy soil in 2-liter pots and irrigated either with ammonium or nitrate solution in concentrations (5 mg l⁻¹ N) commonly found in clearcut boreal forest sites. Irrigation with pure water was used as the control treatment. In a further experiment (Exp. 2), seedlings were irrigated similarly, but irrigation with ammonium and nitrate mixture in proportion of 1:2 (total 5 mg l⁻¹ N) was used as the control treatment. In addition, N treatments were combined with contrasting soil temperatures of 11.5 and 21 °C. Irrigation with N-free pure water resulted in decreased root growth (Exp. 1). In both experiments, no clear differences were found in shoot or root growth in the different N-source treatments. Regardless of N application, the most pronounced effect was increased root growth at the higher soil temperature (Exp. 2). The results question the importance of N form on the growth of Norway spruce seedlings in boreal forests.

Key words: nitrogen, nutrient availability, *Picea abies*, root egress, soil temperature

Introduction

In the boreal zone, Norway spruce (*Picea abies*) is widely planted for reforestation (Grossnickle 2000). In Finland, for example, about 60% of a total of ca. 150 million seedlings planted each year are Norway spruce (Finnish For. Res. Inst. 2002). A couple of years after outplanting to forest sites, stagnation of seedling growth in Norway spruce and other spruce species has been recorded (Björkman 1953, Vyse 1981, Grossnickle 2000). Planting shock is often associated with post-planting water stress, but with

or without water stress, this slow growth may also be related to retarded adaptation to mycorrhizal colonization and to low uptake of nitrogen (N) and other nutrients after planting (Björkman 1953, Grossnickle 2000).

In general, N is the most growth-limiting nutrient in the soil of boreal forests (e.g. Viro 1965). Limited N availability is partly due to low soil temperatures, which decrease water viscosity and mass flow in the soil as well as cell metabolic activity (Lopushinsky & Kaufman 1984, Marschner *et al.* 1991, Ryyppö *et al.* 1998). After clearcutting and site preparation, soil tempera-

ture tends to increase and thus affect nutritional and other growth prerequisites (Örlander *et al.* 1990, Kubin & Kempainen 1994). In addition, N availability to seedlings is affected by soil organic material and its mineralization (Kraske & Fernandez 1990, Nordborg *et al.* 2003). Thus, seedling growth after planting may vary, in addition to seedling material, due to site, method of site preparation and planting spot.

Based mainly on nutrient-uptake studies, Norway spruce has been suggested to prefer ammonium (NH_4^+) to nitrate (NO_3^-) as a source of N (Marschner *et al.* 1991, Buchmann *et al.* 1995, Gessler *et al.* 1998, Högberg *et al.* 1998, George *et al.* 1999, Öhlund & Näsholm 2001). Furthermore, low soil temperatures may limit uptake of nitrate by Norway spruce more than uptake of ammonium (Gessler *et al.* 1998). Organic N can also be utilized by conifer seedlings (Öhlund & Näsholm 2001, Wallander 2002, Persson *et al.* 2003); but recalcitrant organic N may be barely usable (George *et al.* 1999).

When nitrate prevails in the soil, seedlings of late-successional spruces, like Norway spruce, have been suggested to have decreased growth potential relative to other surface vegetation, thus leading to problems in plantation establishment (Kronzucker *et al.* 1997). However, data about the effects of nutrient availability and other soil conditions on growth of planted Norway spruce seedlings is scarce. N uptake by trees has been studied relatively extensively in soilless cultures (e.g. Ingestad 1979, Boxman & Roelofs 1987, Peuke & Tischner 1991, Garnett & Smethurst 1999, Öhlund & Näsholm 2001), but far less in soil (Crabtree & Bazzaz 1993, Buchmann *et al.* 1995, George *et al.* 1999, Persson *et al.* 2003). Furthermore, only a few studies have dealt with the effects of temperature on N availability to any plant species or with the subsequent growth (e.g. Clarkson & Warner 1979, Gessler *et al.* 1998, Garnett & Smethurst 1999, Adam *et al.* 2003).

Thus, there is a need for information about the effects of N source and temperature on the growth of planted Norway spruce seedlings. The aim of the present study was to test whether different combinations of nitrate and ammonium solution and soil temperature affect the growth of transplanted Norway spruce seedlings in sandy soil.

Materials and methods

Preliminary experiment (Exp. 1)

Two-year-old Norway spruce seedlings produced for practical reforestation were used in the experiment. The seedlings were sown using seed-orchard seed of local origin and grown in *Sphagnum* peat in containers (container size 110 cm³, type PL-64F, Lannen Oyj., Iso-Vimma, Finland) according to standard nursery procedures in a greenhouse in central Finland. The seedlings were stored over winter in container trays placed in cardboard boxes in a freezer (−3 °C). Before the experiment, the seedlings were thawed for a week at +8 °C.

A total of 75 vigorous seedlings of uniform height (mean = 28.6, S.D. = 2.5 cm) were selected for the experiment. The roots of the seedlings were gently washed free of the peat medium. Most of the seedlings roots were mycorrhizal (ca. 80%–90%), almost solely with *Laccaria* sp. Fifteen seedlings were measured for height, diameter at root collar, shoot and root volumes (Harrington *et al.* 1994) and biomass fractions (as dry mass at 65 °C). In January 2002, the rest of the seedlings were transplanted into 2-liter pots filled with sand containing 23.4% by mass of fractions 0.2–0.6 mm in diameter and 5.2% of fractions below 0.2 mm in diameter. Soil organic matter concentration was 0.6% by mass. Soil bulk density was ca. 1.50–1.55 g cm⁻³.

The transplanted seedlings were divided into three groups (3 × 20 seedlings), each of which received either ammonium solution, nitrate solution or pure water. Ammonium was applied as NH_4Cl and nitrate as KNO_3 . Both solutions applied had a concentration of 5 mg l⁻¹ N, which is similar to that found on clearcut forest sites (e.g. Nohrstedt *et al.* 1996, Smolander *et al.* 2000). The used solutions were buffered to pH 4.5 with HCl since the most favorable soil pH for Norway spruce and Scots pine (*Pinus sylvestris*) seedlings is 4–5 (Rikala & Jozefek 1990) and nitrification is very limited at pH 4 or lower (Persson & Wiren 1995). Application of pure deionized water, which was also buffered to pH 4.5, was used as the control treatment.

Combined samples within treatments were collected from the leachates from the pots three

times during the experiment. Leachates were analyzed for total N, NH_4 and $(\text{NO}_2 + \text{NO}_3)$ with a flow-injection analyser (FIA) or ion chromatography. Organic N was estimated as total N – NH_4 – $(\text{NO}_2 + \text{NO}_3)$. During the experiment, the total N concentration of the leachates was higher in the KNO_3 treatment (6–9 mg l^{-1}) than in the other treatments (< 3 mg l^{-1}). In the NH_4Cl treatment, the slightly elevated NO_3^- concentrations (< 3 mg l^{-1}) indicated that some nitrification has occurred (< 0.1 mg l^{-1} in H_2O treatment).

Natural light in the greenhouse was supplemented with artificial lighting to give a photoperiod of 18/6 h and photosynthetically active radiation varying spatially from 280 to 330 $\mu\text{mol m}^{-2} \text{s}^{-1}$ at the seedlings' shoot level on cloudy days. On cloudy days, the mean daily and nocturnal air temperatures were 20 and 15 °C, respectively. Relative humidity varied from 30% to 50%. The seedlings were irrigated with the solutions twice per week, which kept the soil adequately moist and the solution available to the seedlings. After two months, the seedlings were harvested and measured for the same attributes as the sample seedlings had been prior to transplanting.

Further experiment (Exp. 2)

A total of 105 vigorous two-year-old seedlings of the same origin as in Exp. 1 with uniform height (mean = 33.9 cm, S.D. = 2.1) were selected for this experiment. Again, 15 seedlings were measured before transplanting and the rest were transplanted as in Exp. 1. Then 90 seedlings ($2 \times 3 \times 15$) were divided into 2×3 treatment groups, in which two soil-temperature and three N-solu-

tion treatments were applied (Table 1). After transplanting, the seedlings were grown from February to April 2003, after which they were harvested and measured.

Two soil temperatures (11.5 and 21 °C) were applied since they were considered to have a contrasting effect on N utilization by the seedlings (Gessler *et al.* 1998). Soil temperatures were adjusted with a water bath, in which the seedling pots were submerged so that their upper rim was ca. 5 cm above the water level. Seedling pots were covered with an insulating polyurethane layer with holes for the seedling shoots. During the experiment, both temperatures were applied simultaneously in two separate basins. Temperatures in the water bath, pots and ambient air were recorded with a datalogger (Table 2). The temperature basins were thus not replicated independently, but the possible dependency of observations was minimized by changing the seedling positions weekly within the temperature blocks and providing as homogenous lighting and seedling irrigation as possible for both basins.

Ammonium and nitrate solutions were applied to both temperature treatments as in Exp. 1. However, a combined solution (5 mg l^{-1} N) of NH_4Cl and KNO_3 in the proportion 1:2 N was used as the control treatment, since similar ammonium-to-nitrate ratios have been found in the field (Smolander *et al.* 2000). In addition, in order to reduce possible nitrification and subsequent formation of NO_3^- , all N solutions were buffered with HCl to pH 4.2. Leachates from the pots were analyzed as in Exp. 1. N concentration in the leachates was higher in the KNO_3 and the combined solution treatment at a soil temperature 21 °C (< 4.5 and < 2.5 mg l^{-1} , respectively) than in the other treatments during the experiment (< 1.5 mg l^{-1}). In the NH_4Cl treat-

Table 1. Solutions applied (5 mg l^{-1} N) and prevailing mean temperature in the water bath in the treatments used in Exp. 2.

Treatment	Solution	Temperature (°C)
Amm_LT	NH_4Cl	11.5
Nit_LT	KNO_3	11.5
Mix_LT	$\text{NH}_4\text{Cl} + \text{KNO}_3$	11.5
Amm_HT	NH_4Cl	21.0
Nit_HT	KNO_3	21.0
Mix_HT	$\text{NH}_4\text{Cl} + \text{KNO}_3$	21.0

Table 2. Temperatures (°C) in the treatments used during Exp. 2.

Position	Treatment	Mean	S.D.
Water bath	Lower temperature	11.5	1.4
Water bath	Higher temperature	20.9	2.0
Pots	Lower temperature	12.5	1.6
Pots	Higher temperature	20.6	1.9
Air	Both temperatures	19.3	2.4

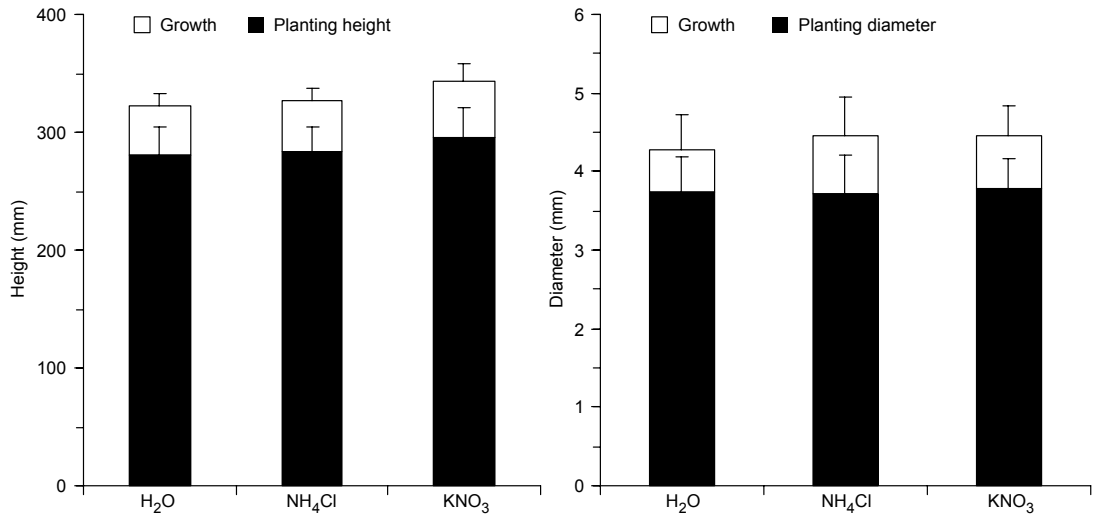


Fig. 1. Seedling growth in height and in diameter at root collar in the treatments used in Exp. 1 (mean + S.D.).

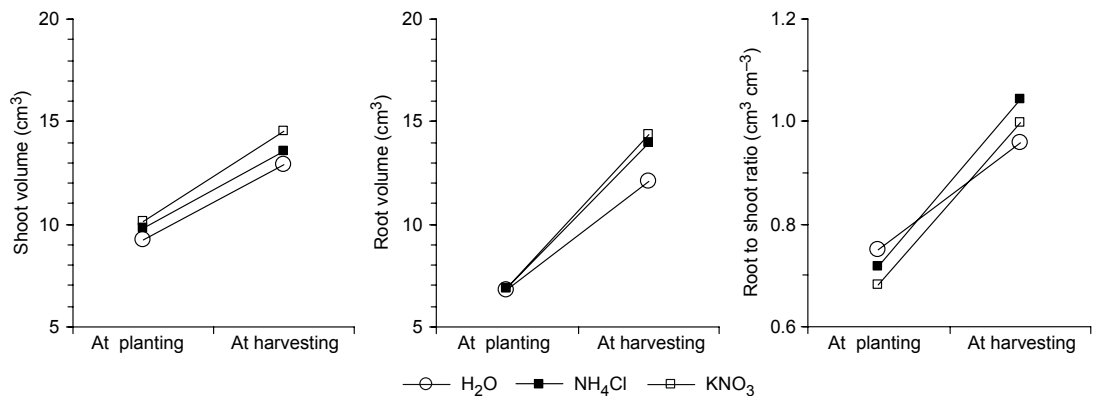


Fig. 2. Mean shoot- and root-volume growths in the treatments used in Exp. 1.

ment, the slightly elevated NO_3^- concentrations ($< 0.4 \text{ mg l}^{-1}$) indicated that little nitrification had occurred.

Data analysis

The seedlings in the three treatments in Exp. 1 were fully randomized, which was ensured by changing the pot positions weekly. Exp. 2 was a randomized block design in which the seedlings and treatments were positioned randomly within the two temperature treatments. Differences between treatments were analyzed with the analysis of variance (ANOVA) and Tukey's test.

Results

Experiment 1

At planting, mean total foliar N was 1.81% (or 18.1 mg g^{-1}). At harvesting, mean total foliar N was 0.82% in the pure water treatment, 0.90% in the NH_4Cl treatment and 0.95% in the KNO_3 treatment. In new needles at harvesting, foliar N was rather uniform in the treatments (mean = 0.83%). Height and diameter growth did not differ significantly between treatments (ANOVA: $P > 0.05$) (Fig. 1). Of the seedling attributes measured, only root-volume growth was significantly different between treatments

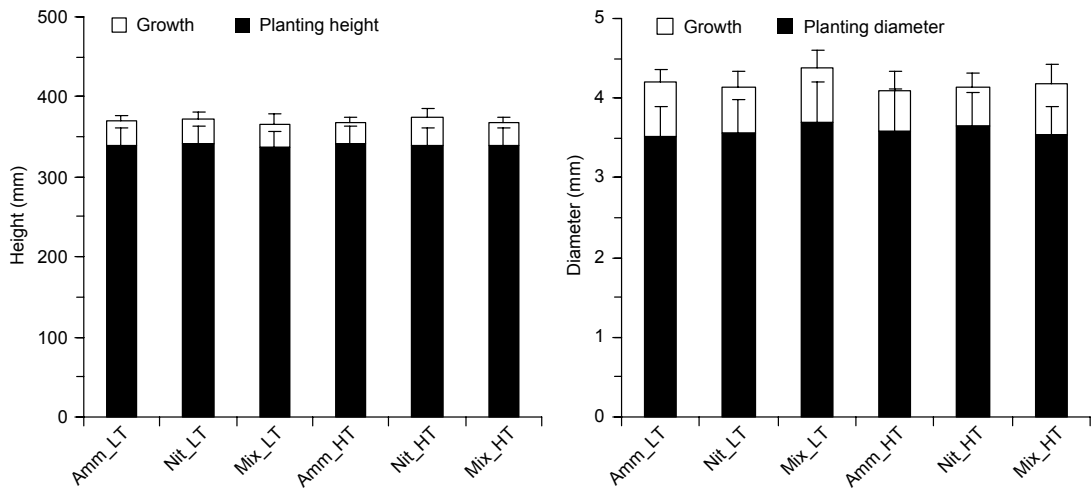


Fig. 3. Seedling growth in height and in diameter at root collar in the treatments used in Exp. 2 (mean + S.D.).

(ANOVA: $P = 0.02$). As compared with N application, irrigation with pure water resulted in decreased root growth (Tukey's test: $P < 0.05$) (Fig. 2).

Experiment 2

Mean total foliar N was 1.27% at planting and 0.46%–0.56% at harvesting (in new needles 0.51%–0.57%). Height growth differed between N-source treatments at $P = 0.061$ (ANOVA) but not between root temperatures (Fig. 3 and Table 3). Diameter growth at the root collar was significantly affected by temperature ($P = 0.009$) and N source applied ($P = 0.041$) but, according to Tukey's test, did not differ between individual treatments ($P > 0.05$). Initial diameter of the root collar at the time of planting was found to be a significant covariate for diameter growth ($P = 0.006$).

Shoot-volume growth did not differ among treatments (Fig. 4 and Table 3). Root-volume growth clearly differed between temperatures (ANOVA: $P < 0.0005$) but not between N-source treatments. The root-to-shoot ratio in terms of volume and dry mass at harvesting also differed significantly between the applied temperatures ($P < 0.0005$) but not between N sources ($P > 0.05$). No other attribute at harvesting differed significantly among treatments.

Discussion

Clearcutting and site preparation tend to increase soil temperature as well as alter moisture conditions and subsequently soil pH and nutrient conditions (Örlander *et al.* 1990, Kubin & Kempainen 1994). It has been suggested that in late-successional forest stands, the soil contains more ammonium than nitrate; while in the early successional stage, the situation is opposite due to increased mineralization and nitrification (Van Cleve *et al.* 1993, Kronzucker *et al.* 1997). The increasing ratio of soil ammonium to nitrate together with differing ammonium transport efficiencies in different tree species may be important forces in competition between species and

Table 3. ANOVA results (P values) for growth of height, diameter at root collar, shoot volume and root volume in Exp. 2. Diameter source denotes the covariate of diameter at planting.

Source	df	P value for growth of			
		height	diameter	shoot vol.	root vol.
Temp.	1	0.451	0.009**	0.487	< 0.0005***
N source	2	0.061	0.041*	0.386	0.787
T × N	2	0.076	0.431	0.673	0.312
Diam.	1		0.006**		
Error	83–84				

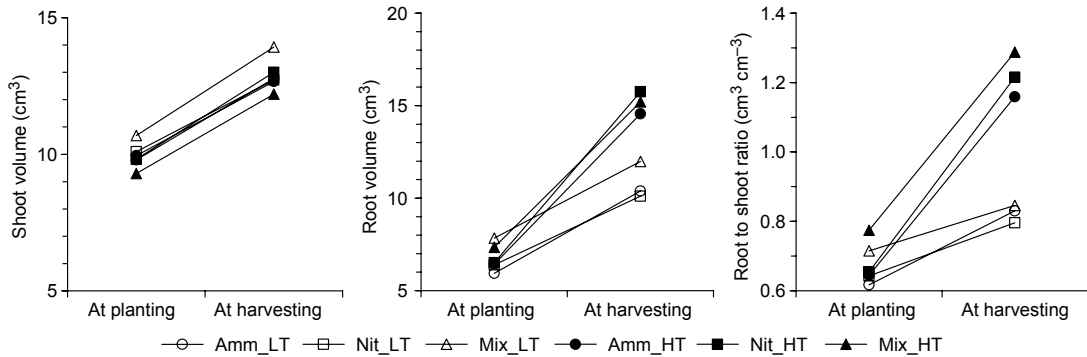


Fig. 4. Mean shoot- and root-volume growths in the treatments used in Exp. 2.

in forest succession (Kronzucker *et al.* 2003). However, little data are available to test this hypothesis.

In practice, clearfelling and site preparation tend to increase the total N concentration in the soil solution (Smolander *et al.* 2000, Högbom *et al.* 2002, Piirainen *et al.* 2002). Furthermore, soil organic N can increase (concentrations below 3.5 mg l^{-1}) (Piirainen *et al.* 2002). In a clearcut Norway spruce stand, ammonium concentrations of ca. 0.5 and 2 mg l^{-1} N have been found in the soil solution in unprepared mineral topsoil and in mounds, respectively (Smolander *et al.* 2000). Corresponding values for nitrate were somewhat higher, ca. 1 and 5 mg l^{-1} N. Similar concentrations have also been found elsewhere (Högberg *et al.* 1986, Termorshuizen & Ket 1991, Nohrstedt *et al.* 1996, Högbom *et al.* 2002). On another clearcut Norway spruce stand (Piirainen *et al.* 2002), somewhat lower nitrate concentrations (below 0.5 mg l^{-1}) were found at locations without site preparation. In areas with higher N deposition, the concentrations of nitrate and ammonium in the soil solution are higher (Gessler *et al.* 1998, cf. Buchmann *et al.* 1995).

In the present study, the N levels available to seedlings were close to natural levels after outplanting to forest sites (e.g. Nohrstedt *et al.* 1996, Smolander *et al.* 2000). With respect to tree-nursery recommendations (Ingestad & Kähr 1985, Landis 1985, Rikala 2002), the foliar N concentrations were relatively low already at the start of the experiments. Therefore, the seedlings had suboptimal N availability as compared with the optimum nutrition; thus showing suboptimal growth rate. However, the use of nutrients in

Finnish tree nurseries has traditionally been low, especially during the second year of seedling growth, since plantation establishment has been found to succeed reasonably well with these N-deficient seedlings (Rikala *et al.* 2004).

With the N concentrations used in this study, the source of inorganic N showed no clear effect on the growth of mycorrhizal Norway spruce seedlings planted in sandy soil at the two contrasting soil temperatures used (11.5 and 21 °C). Regardless of N application, the most pronounced effect was decreased root growth at the lower soil temperature (Exp. 2) and during irrigation with N-free pure water (Exp. 1). The sensitivity of the root-growth response to temperature and N availability is probably due to dependence on current photosynthesis, since shoot growth in conifer seedlings can utilize the previous year's photosynthate reserves, while new root growth is more dependent on current photosynthates (van den Driessche 1987).

In general, high foliar N usually indicates better seedling growth during the second season (Malik & Timmer 1996, Nordborg *et al.* 2003). N uptake of planted Norway spruce seedlings depends on soil organic material, and on its mineralization, and increases with root growth (Kraske & Fernandez 1990, Örlander *et al.* 1990, Nordborg *et al.* 2003). According to the present study, restricted root growth due to limited N availability (Exp. 1) and low soil temperature (Exp. 2) during the first growing season after planting may suggest suboptimal preconditions for shoot growth during the next season. In practice, the use of site-preparation methods such as mounding, which increases soil temperature and

includes humus in the planting spot, is beneficial for reforestation with Norway spruce seedlings (Örlander *et al.* 1990).

Norway spruce has been found to take up more ammonium than nitrate but it has also been found to utilize mainly nitrate, which has better mobility in soil (Marschner *et al.* 1991, Lumme 1994, Buchmann *et al.* 1995, Gessler *et al.* 1998, Högberg *et al.* 1998, George *et al.* 1999, Öhlund & Näsholm 2001). The presence of ammonium may decrease nitrate uptake by Norway spruce seedlings (Gessler *et al.* 1998). In pure nitrate solution, Norway spruce and Scots pine seedlings may show relatively minor growth reduction as compared with that in growing solutions with different nitrate-to-ammonium ratios (Ingestad 1979).

On the other hand, Norway spruce seedlings with natural mycorrhizae have been shown not to differ in their uptake of nitrate and ammonium (Lumme & Smolander 1996) nor in their uptake of N from inorganic and organic sources (Persson *et al.* 2003). In the present study, the seedling growth response suggests that mycorrhizae (*Laccaria* sp.) had no effect or affected the uptake of both N forms by seedlings similarly. In the nursery, *Laccaria* sp. mycorrhizae have previously been reported to retard seedling growth as compared with that of uncolonized control seedlings (Pennanen *et al.* unpubl. data). In general, mycorrhizal colonization of Norway spruce seedlings can cause increased uptake of N and elevated CO₂ assimilation, but uptake of ammonium (not of nitrate) may also result in an increased respiration rate (Eltorp & Marschner 1996, Brunner *et al.* 2000, Jentschke *et al.* 2001).

Prevailing soil temperatures in mature stands have been found to limit uptake of nitrate by Norway spruce more than that of ammonium (Gessler *et al.* 1998). Nitrate uptake by red maple has also been found to decrease with decreasing temperature (34–14 °C) (Adam *et al.* 2003). Also other soil conditions may affect the uptake of N (Stamp *et al.* 1997). With decreasing pH from 5.5 to 2.5, nitrate uptake by Norway spruce seedlings has been shown to decrease sharply (Peuke & Tischner 1991). In addition to low pH, ammonium and Al³⁺ can also decrease nitrate uptake (Peuke & Tischner 1991). With

nitrate fertilization, Fe chelate can improve seedling growth (van den Driessche 1978). In a study with four birch species, only one showed a difference in growth depending on the inorganic N source applied; the effect of which was dependent on the light environment (Crabtree & Bazzaz 1993). In another study, silver birch seedlings differed only slightly in their uptake of both inorganic N sources; but biomass growth was higher with nitrate (Vuorinen *et al.* 1995).

In conclusion, most previous nutrient-uptake studies indicate that Norway spruce seedlings are able to utilize both inorganic N forms but usually prefer ammonium. This study, however, showed no clear effect of applying nitrate and ammonium with levels found in natural boreal conditions on the growth of mycorrhizal Norway spruce seedlings at either the low (11.5 °C) or high (21 °C) soil temperature. This discrepancy may be partly due to the lower, natural N application levels and temperatures used for the mycorrhizal seedlings growing in the soil in the present study, while previous studies have mostly focused on nutrient uptake in soilless cultures in controlled lab conditions. Therefore, the results of this study question the importance of N form on the actual growth of Norway spruce seedlings in boreal forests.

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References

- Adam, M. L., Kelly, J. M., Graves, W. R. & Dixon, P. 2003: Net nitrate uptake by red maple is a function of root-zone temperature. — *J. Pl. Nutr.* 26: 203–222.
- Bassirirad, H., Thomas, R. B., Reynolds, J. F. & Strain, B. R. 1996: Differential responses of root uptake kinetics of NH₄⁺ and NO₃⁻ to enriched atmospheric CO₂ concentration in field-grown loblolly pine. — *Pl. Cell Env.* 19: 367–371.
- Bauer, G. A. & Berntson, G. M. 2001: Ammonium and nitrate acquisition by plants in response to elevated CO₂ concentration: the roles of root physiology and architecture. — *Tree Physiol.* 21: 137–144.
- Björkman, E. 1953: Om orsakerna till granens tillväxts-

- vårigheter efter plantering i nordsvensk skogsmark. — *Norrlands Skogsvårdsförbunds Tidskrift*: 285–316.
- Boxman, A. W. & Roelofs, J. G. M. 1988: Some effects of nitrate versus ammonium nutrition on the nutrient fluxes in *Pinus sylvestris* seedlings. Effects of mycorrhizal infection. — *Can. J. Bot.* 66: 1091–1097.
- Brunner, I., Brodbeck, S. & Genenger, M. 2000: Effects of various nitrogen loads on the nitrate reductase activity in roots and mycorrhizas of Norway spruce seedlings. — *Phyton* 40: 43–48.
- Buchmann, N., Schulze, E.-D. & Gebauer, G. 1995: ¹⁵N-ammonium and ¹⁵N-nitrate uptake of a 15-year-old *Picea abies* plantation. — *Oecologia* 102: 361–370.
- Clarkson, D. T. & Warner, A. J. 1979: Relationships between air temperature and transport of ammonium and nitrate ions by Italian and perennial ryegrass. — *Pl. Physiol.* 64: 557–561.
- Crabtree, R. C. & Bazzaz, F. A. 1993: Seedling response of four birch species to simulated nitrogen deposition: ammonium vs. nitrate. — *Ecological App.* 3: 315–321.
- Eltrop, L. & Marschner, H. 1996: Growth and mineral nutrition of nonmycorrhizal and mycorrhizal Norway spruce (*Picea abies*) seedlings grown in semi-hydroponic sand culture. II. Carbon partitioning in plants supplied with ammonium or nitrate. — *New Phytol.* 133: 479–486.
- Finnish For. Res. Inst. 2002: *Finnish statistical yearbook of forestry*.
- Garnett, T. P. & Smethurst, P. J. 1999: Ammonium and nitrate uptake by *Eucalyptus nitens*: effects of pH and temperature. — *Pl. Soil* 214: 133–140.
- George, E., Stober, C. & Seith, B. 1999: The use of different soil nitrogen sources by young Norway spruce plants. — *Trees* 13: 199–205.
- Gessler, A., Schneider, S., von Sengbusch, D., Weber, P., Hanemann, U., Huber, C., Rothe, A., Kreuzer, K. & Rennenberg, H. 1998: Field and laboratory experiments on net uptake of nitrate and ammonium by the roots of spruce (*Picea abies*) and beech (*Fagus sylvatica*) trees. — *New Phytol.* 138: 275–285.
- Grossnickle, S. C. 2000: *Ecophysiology of northern spruce species: The performance of planted seedlings*. — NRC Res. Press, Ottawa.
- Harrington, J. T., Mexal, J. G. & Fisher, J. T. 1994: Volume displacement provides a quick and accurate way to quantify new root production. — *Tree Pl. Not.* 45: 121–124.
- Högberg, P., Högberg, L. & Schinkel, H. 1998: Nitrogen-related root variables of trees along an N-deposition gradient in Europe. — *Tree Physiol.* 18: 823–828.
- Högberg, P., Granström, A., Johansson, T., Lundmark-Thelin, A. & Näsholm, T. 1986: Plant nitrate reductase activity as an indicator of availability of nitrate in forest soils. — *Can. J. For. Res.* 16: 1165–1169.
- Högberg, L., Nilsson, U. & Örlander, G. 2002: Nitrate dynamics after clear felling monitored by in vivo nitrate reductase activity (NRA) and natural ¹⁵N abundance of *Deschampsia flexuosa* (L.) Trin. — *For. Ecol. Manage.* 160: 273–280.
- Ingestad, T. 1979: Mineral nutrient requirements of *Pinus sylvestris* and *Picea abies* seedlings. — *Physiol. Plant.* 45: 373–380.
- Ingestad, T. & Kähr, M. 1985: Nutrition and growth of coniferous seedlings at varied relative nitrogen addition rate. — *Physiol. Plant.* 65: 109–116.
- Jentschke, G., Godbold, D. L. & Brandes, B. 2001: Nitrogen limitation in mycorrhizal Norway spruce (*Picea abies*) seedlings induced mycelial foraging for ammonium: implications for Ca and Mg uptake. — *Pl. Soil* 234: 109–117.
- Kraske, C. R. & Fernandez, I. J. 1990: Conifer seedling growth response to soil type and selected nitrogen availability indices. — *Soil Sci. Soc. Am. J.* 54: 246–251.
- Kronzucker, H. J., Siddiqi, M. Y. & Glass, A. D. M. 1997: Conifer root discrimination against soil nitrate and the ecology of forest succession. — *Nature* 385: 59–61.
- Kronzucker, H. J., Siddiqi, M. Y., Glass, A. D. M. & Britto, D. T. 2003: Root ammonium transport efficiency as a determinant in forest colonization patterns: a hypothesis. — *Physiol. Plant.* 117: 164–170.
- Kubin, E. & Kempainen, L. 1994: Effect of soil preparation of boreal spruce forest on air and soil temperature conditions in forest regeneration areas. — *Acta For. Fennica* 244.
- Landis, T. D. 1985: Mineral nutrition as an index of seedling quality. — In: Duryea, M. L. (ed.), *Evaluating seedling quality procedures, and predictive abilities of major tests*: 29–48. Forest Research Laboratory, Proceedings of the workshop held October 16–18, 1984. Oregon State Univ., Corvallis.
- Lopushinsky, W. & Kaufmann, M. R. 1984: Effects of cold soil on water relations and spring growth of Douglas fir seedlings. — *For. Sci.* 30: 628–634.
- Lumme, I. 1994: Nitrogen uptake of Norway spruce (*Picea abies* Karst.) seedlings from simulated wet deposition. — *For. Ecol. Manage.* 63: 87–96.
- Lumme, I. & Smolander, A. 1996: Effect of nitrogen deposition level on nitrogen uptake and bud burst in Norway spruce (*Picea abies* Karst.) seedlings and N uptake by soil microflora. — *For. Ecol. Manage.* 89: 197–204.
- Malik, M. & Timmer, V. R. 1996: Growth, nutrient dynamics, and interspecific competition of nutrient-loaded black spruce seedlings on a boreal mixed wood site. — *Can. J. For. Res.* 26: 1651–1659.
- Marschner, H., Häussling, M. & George, E. 1991: Ammonium and nitrate uptake rates and rhizosphere pH in non-mycorrhizal roots of Norway spruce [*Picea abies* (L.) Karst.]. — *Trees* 5: 14–21.
- Nohrstedt, H.-Ö., Sikström, U., Ring, E., Näsholm, T., Högberg, P. & Persson, T. 1996: Nitrate in soil water in three spruce stands in southwest Sweden as related to N-deposition and soil, stand, and foliage properties. — *Can. J. For. Res.* 26: 836–848.
- Nordborg, F., Nilsson, U. & Örlander, G. 2003: Effects of different soil treatments on growth and net nitrogen uptake of newly planted *Picea abies* (L.) Karst. seedlings. — *For. Ecol. Manage.* 180: 571–582.
- Öhlund, J. & Näsholm, T. 2001: Growth of conifer seedlings on organic and inorganic nitrogen sources. — *Tree Physiol.* 21: 1319–1326.
- Örlander, G., Gemmel, P. & Hunt, J. 1990: Site preparation: A Swedish overview. — *FRDA Rep.* 105: 1–61.

- Persson, T. & Wiren, A. 1995: Nitrogen mineralization and potential nitrification at different depths in acid forest soils. — *Pl. Soil* 169: 55–65.
- Persson, J., Högberg, P., Ekblad, A., Högberg, M. N., Nordgren, A. & Näsholm, T. 2003: Nitrogen acquisition from inorganic and organic sources by boreal forest plants in the field. — *Oecologia* 137: 252–257.
- Peuke, A. D. & Tischner, R. 1991: Nitrate uptake and reduction of aseptically cultivated spruce seedlings, *Picea abies* (L.) Karst. — *J. Exp. Bot.* 42: 723–728.
- Piirainen, S., Finér, L., Mannerkoski, H. & Starr, M. 2002: Effects of forest clear-cutting on the carbon and nitrogen fluxes through podzolic soil horizons. — *Pl. Soil* 239: 301–311.
- Rikala, R. 2002: Metsätaimiopas — taimien valinta ja käsittely tarhalta uudistusalueelle. — *Metsäntutkimus. tiedonant.* 881: 1–107.
- Rikala, R. & Jozefek, H. J. 1990: Effect of dolomite lime and wood ash on peat substrate and development of tree seedlings. — *Silva Fennica* 24: 323–334.
- Rikala, R., Heiskanen, J. & Lahti, M. 2004: Autumn fertilization in the nursery affects growth of *Picea abies* container seedlings after transplanting. — *Scand. J. For. Res.* 19: 409–414.
- Ryppö, A., Iivonen, S., Rikala, R., Sutinen, M.-L. & Vapaavuori, E. 1998: Responses of Scots pine seedlings to low root zone temperature in spring. — *Physiol. Plant.* 102: 503–512.
- Smolander, A., Paavolainen, L. & Mälkönen, E. 2000: C and N transformations in forest soil after mounding for regeneration. — *For. Ecol. Manage.* 134: 17–28.
- Stamp, P., Feil, B., Schortemeyer, M. & Richner, W. 1997: Responses of roots to low temperatures and nitrogen forms. — In: Anderson, H. M., Barlow, P. W., Clarkson, D. T., Jackson, M. B. & Shewry, P. R. (eds.), *Plant roots — from cells to systems*: 143–154. Kluwer Acad. Publ., Dordrecht.
- Taylor, C. M. A. & Tabbush, P. M. 1990: Nitrogen deficiency in Sitka spruce plantations. — *For. Comm. Bull.* 89: 20.
- Termorshuizen, A. J. & Ket, P. C. 1991: Effects of ammonium and nitrate on mycorrhizal seedlings of *Pinus sylvestris*. — *Eur. J. For. Pathol.* 21: 404–413.
- Van Cleve, K., Yarie, J., Erickson, R. & Dyrness, C. T. 1993: Nitrogen mineralization and nitrification in successional ecosystems on the Tanana River floodplain, interior Alaska. — *Can. J. For. Res.* 23: 970–978.
- van den Driessche, R. 1978: Response of Douglas fir seedlings to nitrate and ammonium nitrogen sources at different levels of pH and iron supply. — *Pl. Soil* 49: 607–623.
- van den Driessche, R. 1987: Importance of current photosynthate to new root growth in planted conifer seedlings. — *Can. J. For. Res.* 17: 776–782.
- Viro, P. J. 1965: Estimation of the effect of forest fertilization. — *Comm. Inst. For. Fenn.* 59, 3: 1–42.
- Vuorinen, A. H., Rossi, P. & Vapaavuori, E. 1995: Combined effect of inorganic carbon and different nitrogen sources in the growth of media on biomass production and nitrogen uptake in young willow and birch plants. — *J. Pl. Physiol.* 147: 236–242.
- Vyse, A. 1981: Growth of young spruce plantations in interior British Columbia. — *For. Chron.* 57: 174–180.
- Wallander, H. 2002: Utilization of organic nitrogen at two different substrate pH by different ectomycorrhizal fungi growing in symbiosis with *Pinus sylvestris* seedlings. — *Pl. Soil* 243: 23–30.