Effect of nitrogen load on growth and photosynthesis of seedlings of the hybrid larch F₁
(Larix gmelinii var. japonica × L. kaempferi) grown on serpentine soil

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Abstract
We studied the growth and photosynthesis of the hybrid larch F₁ (Larix gmelinii var. japonica × L. kaempferi) grown on serpentine soil, and the effects of soil N load, to determine the performance of this species as reforestation material in serpentine regions. We prepared 16 experimental plots (2 m × 4 m in each), eight on serpentine and eight on brown forest soil, and planted one-year-old cutting seedlings of the hybrid larch F₁ in each plot, in May 2007. Ammonium sulfate was supplied to half of the plots of each soil type in 2008 and 2009, at a load of 47 kg N ha⁻¹ year⁻¹. Although the growth and photosynthetic capacity of hybrid larch F₁ seedlings in the serpentine soil were limited, the rate of growth in serpentine soil was greater than that of Sakhalin spruce (Picea glehnii) that is dominant species in serpentine regions. There was significant interaction between soil type and N load for the growth and photosynthetic parameters. The N load adversely affected growth and photosynthetic parameters in the serpentine soil, while improved them in brown forest soil. Although the growth rate of hybrid larch F₁ without N loading showed high potential as an afforestation species in serpentine region, increasing deposition of N might be a threat to the growth and photosynthesis of the hybrid larch F₁ in serpentine soil.

Key words
gas exchange; infertile soil; nitrogen deposition; nutrient; reforestation material

1. Introduction
Larix species are distributed broadly in cool temperate regions of the northern hemisphere regions (Gower and Richards, 1990). The Japanese larch (Larix kaempferi) was introduced from the central subalpine region of Japan to northern Japan as a trial
plantation species in the 1870s, because it grows faster and has better tolerance to cold than other traditional silvicultural species. Unfortunately it has serious disadvantages as a plantation species there, including susceptibility to shoot blight diseases and grazing by voles (Ito, 1963; Koike et al., 2000, 2004). To overcome these problems, a hybrid larch was developed by crossing *Larix gmelinii var. japonica* and *L. kaempferi*. The resulting hybrid larch F₁ grew faster than the Japanese larch and had a better capacity for carbon accumulation (Miyaki, 1990; Kita et al., 2009).

Serpentine soils are characterized by an excess of elements such as nickel (Ni), chrome (Cr) and magnesium (Mg), and a low ratio of Calcium (Ca) to Mg; these soils are found worldwide (Proctor, 1971; Brooks, 1987; Roberts and Proctor, 1992; Kayama et al., 2009). Many plant species that are intolerant to serpentine soil suffer from toxicity in such regions. Also, low availability of several essential plant nutrients such as phosphorus (P) is a typical trait of serpentine soil (e.g. Kayama et al. 2009). Accordingly, specialized flora grows there, with endemic species distribution (Brooks, 1987; Roberts and Proctor, 1992). If the vegetation of a serpentine region is disturbed, its recovery is very slow (Curtis and Claassen, 2005). Moreover, serpentine regions are often disturbed by erosion and landslides (Lee et al., 2004; O'Dell and Claassen, 2006; Kayama et al., 2007). It is therefore important to identify tree species suited to reforestation of such regions, to prevent disasters and conserve the environment. As larch species can survive in infertile soil, they have been recognized as useful for timber production and absorption of atmospheric CO₂ as a carbon sink, and also afforestation in infertile soil (Zhang et al., 2000; Ryu et al., 2009; Ryu et al., 2010; Watanabe et al., 2011). Kayama et al. (2009) reported that Japanese larch seedlings grew better on serpentine soil than the Sakhalin spruce (*Picea glehnii*), which is a dominant species in serpentine regions of northern Japan.
The atmospheric deposition of nitrogen (N) into terrestrial ecosystems has recently been increasing due to increased anthropogenic emission of N (Galloway et al., 2008; Ohara et al., 2007). Several researchers have reported high N deposition, of 10 to 20 kg N ha\(^{-1}\) year\(^{-1}\) by wet N deposition (bulk precipitation) and 10 to 50 kg N ha\(^{-1}\) year\(^{-1}\) by throughfall and stemflow, in forested areas of Japan (Baba and Okazaki, 1998; Baba et al., 2001; Okochi and Igawa, 2001; Sase et al., 2008; Kimura et al., 2009). In general, N is a limiting resource for tree growth in forest ecosystems, and atmospheric N deposition acts as a fertilizer (e.g. Bobbink et al., 2003; Magnani et al., 2007). However, there are several reports that experimental N supply to soil did not enhance tree growth (Nakaji et al., 2001; Watanabe et al., 2006). The main factor limiting plant growth in serpentine region may be excess Ni and Mg rather than N deficiency, and/or low availability of other nutrients. On this basis, we suggest that an increase in N deposition would not enhance the growth of hybrid larch F\(_1\) in serpentine soil. We tested this hypothesis by studying the effects of N load on the growth and photosynthesis of hybrid larch F\(_1\) seedlings grown on serpentine soil.

2. Materials and methods

2.1. Study site

This study was conducted in the Teshio Experimental Forest of Hokkaido University (45°06’N, 142°12’E, 110 m a.s.l.). Serpentine soils are distributed in the eastern part of the Teshio Experimental Forest, and Sakhalin spruce is dominant in this region (Tatewaki and Igarashi, 1971, Nakata and Kojima, 1987). The mean annual precipitation was approximately 1200 mm. The annual mean, maximum and minimum temperatures were 5.2, 32.0 and −33.5°C, respectively, for the recent 30 years as measured by a thermo-recorder at the Kami-toikan Meteorological Station, about 7 km far from study
We prepared eight experimental plots (each 2 m × 4 m) in each serpentine soil and brown forest soil as controls. According to the FAO-UNESCO systems, serpentine soil and brown forest soil were classified as Podzol and Cambisol, respectively (Nakata and Kojima, 1987). There was a fault between the two soil sites. Climates of the plots were considered to be almost identical because the distances between two soil sites were within 100 m (Kayama et al. 2005). The experimental plots within each soil type were adjoined.

2.2. Plant materials and nitrogen load

In May 2007, cutting seedlings of one-year-old hybrid larch F₁ were planted in experimental plot, four seedlings to each plot and thereby 64 seedlings in total. Each plot was weeded periodically thereafter. After planting, the seedlings were grown for one year without any treatment, to acclimatise to the conditions at the study site.

In May 2008, pellets of ammonium sulfate (commercial fertilizer, Akagi horticulture Co., Ltd., Gunma, Japan) were homogeneously supplied to the soil of four plots in each soil type by handwork. The pellet is usually dissolved with one event of precipitation because of their high solubility. The rate of N load was 47 kg N ha⁻¹ year⁻¹ and this amount is matched with the high N deposition in central Japan (Kimura et al., 2009). We supplied half amount of N (23.5 kg N ha⁻¹) in May 2008 and 2009 for stimulating a seasonal variation of N deposition. Because the Teshio Experimental forest has relatively high amount of snow, which depth in mid winter reaches 100-150 cm, large portion of N is supplied into soil with snow melting in the spring (Nomura et al., 1999; Satoh et al. 1999). The rest N (23.5 kg N ha⁻¹) was supplied in August and November in 2008 and August in 2009.
2.3. Soil properties

In July 2009 we collected a soil core (100 ml) from each plot (four cores per treatment), to evaluate the physical properties of the soil: the dry bulk density, soil water content and maximum water holding capacity. After collection we measured the weight of the soil core \((M_1)\). It was then placed on a tray holding shallow water for 24 hours at room temperature. The resulting saturated soil core was weighed \((M_2)\). It was then dried in an oven for seven days at 105°C, and weighed \((M_3)\). The dry bulk density was calculated as \(M_3\) divided by 100 ml volume of soil cores \((\text{g cm}^{-3})\). The soil water content and maximum water holding capacity were calculated as follows:

Soil water content \((\text{g g}^{-1})\) = \(\frac{M_1 - M_3}{M_3}\)

Maximum water holding capacity \((\text{g g}^{-1})\) = \(\frac{M_2 - M_3}{M_3}\)

To determine the chemical properties of the soil, including pH, total carbon (C) and N concentrations, inorganic N (i.e. NO\(_3^-\) and NH\(_4^+\)), available P, exchangeable cation, and acid soluble metals, we collected soil samples at the end of the experiment in September 2009. Soil samples were collected from each plot (four samples per treatment), and were air-dried for one month. The dried soil samples were crushed with a mortar and passed through a sieve of mesh 2 \(\times\) 2 mm. This soil sample was used for the following measurements.

The soil pH was measured with a pH meter (HM-30V, DKK-TOA Co. Ltd., Tokyo, Japan) after 10 g soil sample had been shaken with 50 ml of ion-exchanged water for 1 h (van Reeuwijk, 1993). The total C and N concentrations in the soil was measured
by gas chromatography (GC-8A, Shimadzu, Kyoto, Japan) after combustion with circulating O\textsubscript{2} by an NC analyzer (Sumigraph NC-900, Sumika Chemical Analysis Service, Osaka, Japan). Available P was extracted by the Bray-2 method (Bray and Kurtz, 1945), as follows: 1 g soil sample was shaken with extracting solution containing 0.03 M ammonium fluoride and 0.1 M hydrochloric acid. The P extracted was determined by the molybdenum blue method with ascorbic acid, using a spectrophotometer (Gene spec III, Hitachi, Tokyo, Japan) (Murphy and Riley, 1962). To extract exchangeable cations from the soil, 5 g soil sample was mixed with 50 ml of 1 M ammonium acetate solution and was shaken for 1 h (Thomas, 1982). Solutions for analyzing acid soluble metals were obtained by adding 50 ml 0.1 M hydrochloric acid to 5 g soil sample with shaking for 1 h (Baker and Amacher, 1982). The concentrations of elements in the sample solutions were determined using an inductively coupled plasma optical emission spectrometer (ICP-OES, IRIS, Jarrel Ash, Franklin, MA, USA). We determined the concentrations of Ca, Mg, potassium (K), Cr and Ni as exchangeable cations, and Cr and Ni as an acid-soluble metal.

2.4. Growth measurement

In May and October 2008 and in May, July and September 2009, we measured the height and stem diameter of all seedlings. In September 2009, all seedlings were harvested in order to determine the dry mass of the plant organs. The seedlings were separated into needles, stems and roots. The plant organs were dried at 70˚C for 1 week and weighed.

2.5. Measurement of gas exchange rates in needles

The gas exchange rates of fully expanded young needles were measured in July 2009 using an open gas exchange system (LI-6400, Li-Cor Inc., Lincoln, NE, USA). Two
seedlings per treatment-plot combination were selected randomly for measurement, to a total of eight seedlings per treatment. The gas exchange rates were measured between 0800 and 1500 hours. The needle temperature during the measurement was maintained at 25 ± 0.5°C and the photosynthetic photon flux density at 1500 μmol m⁻² s⁻¹. The vapor pressure deficit was 1.2 ± 0.2 kPa. To obtain the intercellular CO₂ concentration (Cᵢ)-response curve of the net photosynthetic rate (A), i.e., the A/Cᵢ curve, A was determined at ten CO₂ concentration levels in the chamber (Cₐ, 60-1700 μmol mol⁻¹). We determined A, the transpiration rate and stomatal conductance at 380 μmol mol⁻¹ CO₂ (A₃₈₀, E and Gₛ, respectively), and A at 1700 μmol mol⁻¹ CO₂ (Aₘₐₓ). The stomatal limitation of photosynthesis, maximum rate of carboxylation (Vₖₐₓ) and the maximum rate of electron transport (Jₘₐₓ) were calculated from the A/Cᵢ curve (Farquhar et al., 1980; Long and Bernacchi, 2003). Values of the Rubisco Michaelis constants for CO₂ (Kᵣ) and O₂ (Kₒ) and CO₂ compensation point in the absence of dark respiration (Γ*), for analysis of the A/Cᵢ curve, were as according to Bernacchi et al. (2001) All gas exchange parameters were expressed on the basis of the projected needle area, which was measured with an image scanner (CanoScan LiDE 600F, Cannon, Tokyo, Japan).

2.6. Measurement of needle traits

After the gas exchange rate had been measured, the needles were collected in order to determine the leaf mass per area (LMA) and the chlorophyll content and amount of nutrient elements. After the needle projected area was measured using an image scanner, the samples for chlorophyll quantification were frozen and stored in a freezer at -80°C. Chlorophyll in needles was extracted with dimethyl sulfoxide as according to Barnes et al. (1992) and Shinano et al. (1996), and was determined using a spectrophotometer (Gene spec III, Hitachi, Tokyo, Japan). The other samples, for determination of LMA and
nutrient elements, were dried in an oven for five days at 70°C. The LMA was calculated as the ratio of dry mass to the area of the needles. The dried needles were ground to a fine powder with a sample mill. The N content of the needle sample was determined with a gas chromatography-NC analyzer system, as mentioned above. We calculated the ratio of \( A_{380} \) to the N content as photosynthetic N use efficiency (PNUE). To measure the concentrations of P, Ca, Mg, K, and Ni in the needles, the powdered sample was digested with nitric acid, hydrochloric acid, and hydrogen peroxide. The concentrations of elements in the sample solutions were determined using an ICP-OES (ICPS-7000, Shimadzu, Kyoto, Japan).

2.7. Estimation of nitrogen allocation to photosynthetic functions

The photosynthetic apparatus was divided into three parts: Rubisco, bioenergetics (electron carriers except for photosystems, coupling factor and Calvin cycle enzymes except Rubisco), and light-harvesting complex and photosystems; the fraction of needle N deployed in each function is denoted by N1, N2 and N3. We estimated N1 according to the following equation (Niinemets et al., 1999; Tissue and Lewis, 2010):

\[
N1 = \frac{V_{\text{max}}}{(6.25 \cdot V_{\text{cr}} \cdot \text{LMA} \cdot N_M)}
\]

where \( V_{cr} \) is the specific activity of Rubisco (the maximum rate of RuBP carboxylation per unit Rubisco protein), \( N_M \) denotes N per unit leaf mass, and the factor of 6.25 (g Rubisco [g N in Rubisco]^{-1}) converts N content to protein content. Here, \( V_{cr} \) is equal to 20.5 \( \mu \text{mol} \text{ CO}_2 \cdot (\text{g Rubisco})^{-1} \cdot \text{s}^{-1} \) at 25 °C for purified Rubisco enzyme from Spinacia oleracea (Jordan and Ogren, 1984). Because this method cannot determine the amount of inactivated Rubisco, the calculated value of N1 is an underestimate (Warren and Adams,
The N2 was estimated from gas exchange characteristics according to the equation (Kitaoka and Koike, 2004; Takashima et al., 2004):

\[ N2 = \frac{J_{\text{max}}}{156 \times 9.53 \text{ LMA } N_m} \]

We assume that N in bioenergetics is proportional to \( J_{\text{max}} \), where the ratio of \( J_{\text{max}} \) to the cytochrome f content is 156 mmol mol\(^{-1}\) s\(^{-1}\) (Niinemets and Tenhunen, 1997), and N in bioenergetics per unit cytochrome f is 9.53 mol mmol\(^{-1}\) (Hikosaka and Terashima, 1995). N3 was estimated according to the following equation.

\[ N3 = \frac{37.1 \text{ Chl}}{(\text{LMA } N_m)} \]

where Chl is chlorophyll content (mol m\(^{-2}\)) and the N content per unit chlorophyll is 37.1 mol mol\(^{-1}\) (Evans and Seemann, 1989).

### 2.8. Statistical analysis

Statistical analyses were undertaken using R software, version 2.8.1 (R Development Core Team, 2009). Student's t-test was applied to the difference in physical properties between the two soil types. Two-way analysis of variance (ANOVA) was used to test the effects of soil type and N treatment. In the analysis, a plot was nested within each soil and N treatment and was added to the model as a random effect (Underwood, 1981). When a significant interaction between soil type and N treatment was detected, Fischer's least significant difference test was performed to identify significant differences between the four treatments.
3. Results

3.1. Soil properties

The soil water content and maximum water holding capacity of serpentine soil were significantly higher than values in brown forest soil, but there was no significant difference in dry bulk density between the soil types (Table 1).

The content of Ca, Mg, acid-extracted Cr and Ni (in both extractions) in serpentine soil was each significantly higher than in brown forest soil, whereas the P content was significantly lower in serpentine soil (Table 2). The N load caused a significant increase in the total N and NO$_3^-$ content, and a fall in the K content. Significant interactions between soil type and N load were observed in the values of the pH, the NH$_4^+$ content and the ammonium acetate-extracted Ni content. The pH was higher in serpentine soil than in brown forest soil, and decreased with the N load. This tendency in pH did not change despite the significant interaction. An increase in the content of NH$_4^+$ and Ni induced by N loading was found in serpentine soil, although N loading did not affect the Ni content of brown forest soil.

3.2. Growth response

Figure 1 shows trends in the increments of height and stem diameter of the seedlings. For serpentine soil, all parameters were lower than those in brown forest soil. N loading tended to increase growth parameters in brown forest soil but reduce them in serpentine soil. The dry mass of all organs and the whole-plant dry mass of the seedlings in serpentine soil were significantly lower than those in brown forest soil (Table 3). There was significant interaction between soil type and N load for the dry mass of the stem and roots, and also the whole-plant dry mass. N loading did not significantly affect (rather decrease) the dry mass of seedlings grown on serpentine soil, whereas in brown forest
soil N it led to a significant increase in the root, and a marginally significant increase in stem and whole-plant dry mass. (The $P$ values were 0.074 for root and 0.100 for whole-plant dry mass.)

3.3. Needle gas exchange parameters and traits

All parameters except for stomatal limitation and the LMA of the needles in serpentine soil were significantly lower than the corresponding values in brown forest soil (Table 4). Stomatal limitation and LMA responded oppositely from the other parameters to soil type. The N load significantly increased the value of $N_{\text{area}}$. For $A_{\text{max}}$, $E$, $V_{\text{cmax}}$, $J_{\text{max}}$ and PNUE there was significant interaction between soil type and N loading. The N load significantly reduced the values of these parameters in serpentine soil, but no significant effect of N loading was found in brown forest soil. The interaction of soil type and N loading for $A_{380}$ was marginally significant ($P = 0.058$) and showed a similar trend. N allocation to photosynthetic functions in needles is shown in Figure 2. N loading significantly decreases N allocation to photosynthetic apparatus in serpentine soil but not in brown forest soil. A similar trend in N allocation was observed in each component of photosynthetic function (i.e. N1, N2 and N3).

3.4. Concentrations of elements in needles

Mass-based N, P, K and Ni concentrations in needles of seedlings grown in serpentine soil were significantly lower than those in brown forest soil, but the Mg concentration was higher for serpentine soil (Fig. 3). N loading significantly increased the mass-based N concentration. There was no significant interaction between soil type and N loading for the concentration of any element in needles. The concentration of Cr in needles was quite low and we could not detect Cr in about half of the samples. Figure 4 shows the relations
between $A_{380}$ and amounts of elements in needles. Significant positive correlations were observed between $A_{380}$ and the N, P, K and Ni content in needles, but $A_{380}$ was negatively correlated with Mg content in the needles.

4. Discussion

4.1. Growth and photosynthesis of hybrid larch $F_1$ seedlings grown in serpentine soil

In serpentine soil, the growth and photosynthetic capacity of hybrid larch $F_1$ seedlings were severely limited compared to those in brown forest soil (Fig. 1 and Table 3). High soil water content, high concentrations of Ni and Mg, and a low concentration of P was found for the serpentine soil in the present study site (Table 1, Table 2). Although the decrease in the T/R ratio is considered to be an accommodation to the low P availability (Lambers et al., 2008), the P concentration in needles on serpentine soil was lower than on brown forest soil (Table 3, Fig. 3). Needle N and K concentrations were also lower in serpentine soil, even there was no significant difference between the two soil types in the availability of those elements. Furthermore, although Ca availability was higher in serpentine soil, there was no difference in the concentration of Ca in needles between two soil types. Excess Mg availability in serpentine soil might inhibit the uptake of the other nutrient by root or transport from root to above ground as indicated by Kobayashi et al. (2005). Accommodation to serpentine soil is therefore insufficient to compensate for its infertile condition.

Photosynthetic activity is generally regulated by the foliar N content (Lambers et al., 2008). Lower N content was observed in the needles of seedlings grown on serpentine soil, and this is believed to be the factor responsible for decreasing the photosynthetic activity (Table 4, Fig. 4). The PNUE also decreased in serpentine soil. The decrease in the CO$_2$ uptake owing to stomatal closure led to the decrease in PNUE (Table
4). The content of other elements than N in needles may also affect PNUE (Watanabe et al., 2011). Pâques (1994) reports that the optimal concentrations of elements for growth in needles of the hybrid larch (Larix × eurolepis Henry) growth are 19.5-27.3 mg g⁻¹ for N, 1.4-2.4 mg g⁻¹ for P, 5.2-8.6 mg g⁻¹ for K, 3.3-6.2 mg g⁻¹ for Ca and 1.1-2.0 mg g⁻¹ for Mg. We found that the concentrations of N, P and Ca in needles grown on serpentine soil were less than the optimum (Fig. 3). The value of $A_{380}$ was significantly positively correlated with the N and P concentrations (Fig. 4). The concentration of P in needles therefore appears to be a further factor that decreases photosynthetic activity in serpentine soil, consistent with Kayama et al. (2009). This possibility can be verified by an experiment of P fertilization in the future. The concentration of Mg in needles was significantly negatively correlated with $A_{380}$ (Fig. 4). However, the needle Mg concentrations on serpentine soil were not high enough to decrease photosynthetic activity and growth (Rao et al., 1987; Kobayashi et al., 2005).

Surprisingly, the Ni concentration in the needles of seedlings grown on serpentine soil was lower than that for brown forest soil (Fig. 3). Ectomycorrhizal symbiosis plays an important role in the exclusion of Ni (Hartley et al., 1997; Leyval et al., 1997; Kayama et al., 2005, 2006) and we observed the formation of ectomycorrhizal roots in all seedlings in the present study (data not shown). In general, infection level to ectomycorrhiza is higher in infertile soil than that in fertile soil (van der Heijden and Sanders, 2002). Although we did not quantitatively evaluate the infection level to ectomycorrhiza in root, the extent of the infection level in serpentine soil may be higher than that in brown forest soil. Actually, the relative C allocation to roots was estimated to be higher in serpentine soil, because of the lower T/R ratio (Table 3), and that to ectomycorrhiza may therefore be higher, giving rise to well-developed ectomycorrhiza which robustly exclude Ni in serpentine soil. On the other hand, the species composition
of ectomycorrhiza is quite dependent on the soil type (van der Heijden and Sanders, 2002; Qu et al., 2004). The difference in species composition of ectomycorrhiza between two soil types might also explain the lower absorption of Ni in serpentine soil as compared to that in brown forest soil. As another possibility, greater accumulation of Ni in the needles in brown forest soil may be responsible for the higher transpiration rate (Table 4).

According to Kayama et al. (2005; 2006; 2009), Ni concentration in the matured needle of the seedlings of conifer species (Sakhalin spruce, Yezo spruce (Picea jezoensis Carr.) and Norway spruce (Picea abies Karst) and Japanese larch) grown on serpentine soil were ranged from 15 to 70 μg g⁻¹. Kayama et al. (2009) demonstrated no decline of net photosynthetic rate in the range below 30 μg g⁻¹ of Ni concentration in needles of Japanese larch seedlings grown in the same experimental forest with the present study. The Ni concentration in the needle of hybrid larch F₁ in the present study was rather low (15-25 μg g⁻¹, Fig. 3). Therefore, we consider the Ni concentration in needles was within a range that does not exert a negative effect on photosynthesis, and the correlation between \(A_{380}\) and Ni concentration simply was a result with the other mechanism.

4.2. Effects of N load on the growth and photosynthesis of hybrid larch F₁ seedlings

The N loading increased the growth of the hybrid larch F₁ seedlings grown on brown forest soil, although the increase was only just statistically significant (Table 3). In serpentine soil, we found the opposite: N loading reduced growth. Similar impairment was observed in \(A_{380}\), \(A_{\text{max}}\), \(E\), \(V_{\text{cmax}}\), \(J_{\text{max}}\) and PNUE in the needles (Table 4). The different responses of growth in serpentine soil and brown forest soil to N loading are due to the differing responses in photosynthetic activity. The negative effects of N loading on photosynthetic activity in serpentine soil was mainly due to the reduction in PNUE of the
needles as a result of the decreased nitrogen allocation to photosynthetic apparatus (Table 4, Fig. 2). The concentrations of elements in needles were not the causal factor responsible (Fig. 3). Stomatal limitation may contribute to the response of PNUE to N loading in serpentine soil, although the interaction between soil type and N loading for PNUE was not significant because of the relatively large variations (Table 4). Further study is needed to clarify the effects of N deposition on photosynthesis of the hybrid larch F₁ grown on serpentine soil.

Soil N loading induced a significant increase in the available Ni concentration in soil (Table 2). Ni availability increases with decreasing soil pH (Mizuno, 1967; Seregin and Kozhevnikova, 2006). For N loading of soil we applied ammonium sulfate, so that some of the $\text{NH}_4^+$ should change to $\text{NO}_3^-$ through nitrification processes and release $\text{H}^+$ to the soil (i.e. decrease the pH) (Fisher and Binkley, 2000). Although the available Ni concentration in serpentine soil was significantly increased by N loading, the concentration of Ni in needles was not increased (Table 2, Fig. 3). This might be because ectomycorrhizal fungi act to exclude Ni, as stated above. However, it is possible that the cost of C needed to maintain the ectomycorrhizal symbiosis that prevents seedlings from reflecting the higher Ni concentration is greater in the seedlings grown on serpentine soil with N loading, causing the reduction in growth.

Growth of the hybrid larch F₁ seedlings without N loading is higher than that of the Sakhalin spruce, native species of serpentine regions, and similar to that of seedlings of the Japanese larch (Kayama et al., 2006; 2009). We therefore assert that the hybrid larch F₁ is useful for reforestation in serpentine regions, as well as the Japanese larch. However, the N load-induced reduction in growth and photosynthetic activity on serpentine soil is of concern for reforestation with the hybrid larch F₁ given increasing N deposition.
5. Conclusion

The results of the present study clearly support our hypothesis. N loading adversely affected growth and gas exchange parameters of seedlings grown in serpentine soil, whereas the opposite trends were observed in brown forest soil. The growth rate of hybrid larch F₁ without N loading showed high potential as an afforestation species in serpentine region. However, increasing deposition of N might be a threat to the growth and photosynthesis of the hybrid larch F₁. The emission control for anthropogenic N compound especially in East Asia region is highly important for the afforestation with hybrid larch F₁ in serpentine regions of Japan. In the next step, we should develop more long-term study from seedlings to mature tree to confirm the ability of hybrid larch F₁ as afforestation species with special attention to the increase of N deposition.

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Reference


**Figure captions**

**Fig. 1** Increments of height and stem diameter of hybrid larch F$_1$ seedlings from May 2008 to September 2009. The seedlings were grown on serpentine soil (S) or brown forest soil (B) in combination with N loading at 0 (-N) or 47 kg N ha$^{-1}$ year$^{-1}$ (+N) (n = 16).

**Fig. 2** Nitrogen allocation in the needles of hybrid larch F$_1$ seedlings. The seedlings were grown on serpentine soil (S) or brown forest soil (B) in combination with N loading at 0 (-N) or 47 kg N ha$^{-1}$ year$^{-1}$ (+N) (n = 8). N1, nitrogen allocated to Rubisco; N2, nitrogen allocated to electron carriers except for photosystems, coupling factor and Calvin cycle enzymes apart from Rubisco; N3, nitrogen allocated to light-harvesting complex and photosystems. Values with different letters attached differ significantly for the sum of N1, N2 and N3, with $P < 0.05$.

**Fig. 3** Concentrations of elements in needles of hybrid larch F$_1$ seedlings grown on serpentine soil (S) or brown forest soil (B) in combination with N loading at 0 (-N) or 47 kg N ha$^{-1}$ year$^{-1}$ (+N). The error bar shows SE (n = 8). ANOVA: ** $P < 0.01$; *** $P < 0.001$; n.s. not significant. There was no significant interaction between soil type and N load for any parameter.

**Fig. 4** Relations between mass-based element content and net photosynthetic rate at 380 μmol mol$^{-1}$ CO$_2$ ($A_{380}$) in the needles of hybrid larch F$_1$ seedlings. The seedlings were grown on serpentine soil (S) or brown forest soil (B) in combination with N loading at 0 (-N) or 47 kg N ha$^{-1}$ year$^{-1}$ (+N). Pearson’s correlation test: ** $P < 0.01$; *** $P < 0.001$; n.s. not significant.
Fig. 1

Graph showing the relationship between height increment (cm) and diameter increment (mm) from 2008 to 2009. The graphs compare height increment and diameter increment across different months (M = March, J = June, A = August, S = September, and O = October). The data points are categorized into S-N, S+N, B-N, and B+N for both height and diameter increments.

- **Height increment (cm)**
  - 2008: Small increments ranging from 0 to 50 cm.
  - 2009: Larger increments ranging from 40 to 80 cm.

- **Diameter increment (mm)**
  - 2008: Moderate increments ranging from 0 to 15 mm.
  - 2009: Slightly higher increments ranging from 15 to 30 mm.

The graph highlights the growth patterns and differences between the months and the two years.
Fig. 2
| Element | Soil N | Soil N.s. | Soil N || Soil N.s. |
|---------|--------|-----------|--------|--------|
| N (mg g⁻¹) | *** | N ** | N.s. | N.s. |
| Ca (mg g⁻¹) | *** | N.s. | N.s. | N.s. |
| P (mg g⁻¹) | *** | N. n.s. | N.s. | N.s. |
| K (mg g⁻¹) | *** | N.s. | N.s. | N.s. |
| Mg (mg g⁻¹) | *** | N.s. | N.s. | N.s. |
| Ni (μg g⁻¹) | *** | N.s. | N.s. | N.s. |
Fig. 4

\[ r = 0.724^{**} \]

\[ r = 0.643^{**} \]

\[ r = 0.261^{**} \]

\[ r = -0.747^{**} \]

\[ r = 0.476^{**} \]

\[ r = 0.643^{**} \]
Table 1 Physical properties of serpentine soil and brown forest soil

<table>
<thead>
<tr>
<th></th>
<th>Serpentine soil</th>
<th>Brown forest soil</th>
<th>t-test</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dry bulk density (Mg m⁻³)</td>
<td>0.85 (0.02)</td>
<td>0.87 (0.02)</td>
<td>n.s.</td>
</tr>
<tr>
<td>Soil water content (g g⁻¹)</td>
<td>0.61 (0.03)</td>
<td>0.42 (0.01)</td>
<td>***</td>
</tr>
<tr>
<td>Maximum water holding capacity (g g⁻¹)</td>
<td>0.57 (0.01)</td>
<td>0.51 (0.01)</td>
<td>**</td>
</tr>
</tbody>
</table>

Each value is the mean of 8 determinations; the standard error is shown in parentheses
*t-test: ** $P < 0.01$; *** $P < 0.001$; n.s. not significant
<table>
<thead>
<tr>
<th>Parameter</th>
<th>Serpentine soil -N</th>
<th>Brown forest soil -N</th>
<th>Two-way ANOVA Soil N Soil×N</th>
</tr>
</thead>
<tbody>
<tr>
<td>pH</td>
<td>6.17 (0.04) a</td>
<td>5.86 (0.11) b</td>
<td>5.54 (0.07) c 4.84 (0.10) d</td>
</tr>
<tr>
<td>Total C (%)</td>
<td>2.21 (0.55)</td>
<td>2.10 (0.23)</td>
<td>2.94 (0.33) 2.50 (0.17)</td>
</tr>
<tr>
<td>Total N (%)</td>
<td>0.11 (0.01)</td>
<td>0.17 (0.02)</td>
<td>0.14 (0.02) 0.17 (0.01)</td>
</tr>
<tr>
<td>NO$_3$ (mg kg$^{-1}$)</td>
<td>3.96 (0.72)</td>
<td>9.91 (1.66)</td>
<td>3.82 (0.70) 6.01 (1.63)</td>
</tr>
<tr>
<td>NH$_4^+$ (mg kg$^{-1}$)</td>
<td>6.19 (0.35) b</td>
<td>10.16 (1.30) a</td>
<td>8.21 (0.50) ab 8.61 (0.57) a</td>
</tr>
<tr>
<td>P (mg kg$^{-1}$)</td>
<td>6.01 (1.76)</td>
<td>3.67 (0.73)</td>
<td>11.43 (1.75) 13.94 (2.37)</td>
</tr>
<tr>
<td>Ca (mg kg$^{-1}$)</td>
<td>686 (38)</td>
<td>758 (53)</td>
<td>304 (75) 339 (85)</td>
</tr>
<tr>
<td>Mg (mg kg$^{-1}$)</td>
<td>1490 (32)</td>
<td>1350 (195)</td>
<td>51 (10) 12 (2)</td>
</tr>
<tr>
<td>K (mg kg$^{-1}$)</td>
<td>113 (17)</td>
<td>103 (9)</td>
<td>114 (15) 67 (5)</td>
</tr>
<tr>
<td>Cr (AA) (mg kg$^{-1}$)</td>
<td>2.25 (0.28)</td>
<td>3.48 (0.48)</td>
<td>3.07 (0.49) 2.47 (0.26)</td>
</tr>
<tr>
<td>Ni (AA) (mg kg$^{-1}$)</td>
<td>3.98 (0.58) b</td>
<td>6.96 (0.38) a</td>
<td>0.06 (0.01) c 0.05 (0.01) c</td>
</tr>
<tr>
<td>Cr (HA) (mg kg$^{-1}$)</td>
<td>2.04 (0.55)</td>
<td>1.37 (0.62)</td>
<td>0.49 (0.04) 0.57 (0.07)</td>
</tr>
<tr>
<td>Ni (HA) (mg kg$^{-1}$)</td>
<td>39.11 (1.99)</td>
<td>40.08 (4.12)</td>
<td>0.19 (0.08) 0.13 (0.06)</td>
</tr>
</tbody>
</table>

Cr and Ni were extracted by two solutions, ammonium acetate (AA) and hydrochloric acid (HA). Each value is the mean of four measurements; the standard error is shown in parentheses.

ANOVA: * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$; n.s. not significant

For each parameter, values with different letters attached differ significantly, with $P < 0.05$.
Table 3 Effect of nitrogen loading on the dry mass of plant organs and the ratio of aboveground dry mass to belowground dry mass (T/R) of hybrid larch F₁ seedlings grown on serpentine soil and brown forest soil

<table>
<thead>
<tr>
<th></th>
<th>Serpentine soil</th>
<th>Brown forest soil</th>
<th>Two-way ANOVA</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>-N</td>
<td>+N</td>
<td>-N</td>
</tr>
<tr>
<td>Needle</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>9.5 (2.0)</td>
<td>3.0 (0.7)</td>
<td>36.2 (4.0)</td>
</tr>
<tr>
<td>Stem</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>10.6 (1.9)</td>
<td>4.3 (0.7)</td>
<td>b 52.7 (6.9)</td>
</tr>
<tr>
<td>Root</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>6.9 (1.2)</td>
<td>3.2 (0.5)</td>
<td>c 23.8 (3.7)</td>
</tr>
<tr>
<td>Whole-plant</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>27.0 (5.0)</td>
<td>10.5 (1.9)</td>
<td>b 112.6 (14.3)</td>
</tr>
<tr>
<td>T/R</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>2.5 (0.4)</td>
<td>2.1 (0.3)</td>
<td>3.7 (0.3)</td>
</tr>
</tbody>
</table>

Each value is the mean of 16 measurements; the standard error is shown in parentheses
ANOVA: * P < 0.05; *** P < 0.001; n.s. not significant
For each parameter, values with different letters attached differ significantly, with P < 0.05
Table 4 Effect of nitrogen loading on the needle traits of hybrid larch F1 seedlings grown on serpentine soil and brown forest soil

<table>
<thead>
<tr>
<th></th>
<th>Serpentine soil</th>
<th>Brown forest soil</th>
<th>Two-way ANOVA</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>-N</td>
<td>+N</td>
<td>-N</td>
</tr>
<tr>
<td>$A_{380}$ (μmol m$^{-2}$ s$^{-1}$)</td>
<td>3.9 (0.7)</td>
<td>2.4 (0.5)</td>
<td>7.3 (0.7)</td>
</tr>
<tr>
<td>$A_{max}$ (μmol m$^{-2}$ s$^{-1}$)</td>
<td>16.3 (2.4) b 10.3 (1.3) c</td>
<td>19.6 (2.0) ab 22.0 (1.4) a</td>
<td>0.07 (0.01) 0.04 (0.01)</td>
</tr>
<tr>
<td>$G_s$ (mol m$^{-2}$ s$^{-1}$)</td>
<td>0.07 (0.01)</td>
<td>0.04 (0.01)</td>
<td>1.02 (0.13) b 0.63 (0.09) c</td>
</tr>
<tr>
<td>$Ls$</td>
<td>0.44 (0.03)</td>
<td>0.49 (0.05)</td>
<td>0.44 (0.03)</td>
</tr>
<tr>
<td>$V_{cmax}$ (μmol m$^{-2}$ s$^{-1}$)</td>
<td>51 (6) b 36 (5) c</td>
<td>56 (4) ab 67 (4) a</td>
<td>51 (6) b 36 (5) c</td>
</tr>
<tr>
<td>$J_{max}$ (μmol m$^{-2}$ s$^{-1}$)</td>
<td>127 (14.6) b 85 (10.7) c</td>
<td>127 (9.7) ab 147 (10.5) a</td>
<td>127 (14.6) b 85 (10.7) c</td>
</tr>
<tr>
<td>PNUE (μmol mol$^{-1}$ s$^{-1}$)</td>
<td>48 (7) b 22 (3) c</td>
<td>61 (6) ab 63 (7) a</td>
<td>48 (7) b 22 (3) c</td>
</tr>
<tr>
<td>LMA (g m$^{-2}$)</td>
<td>83.4 (1.5)</td>
<td>84.1 (2.0)</td>
<td>75.7 (1.2)</td>
</tr>
<tr>
<td>$N_{area}$ (g m$^{-2}$)</td>
<td>1.04 (0.08)</td>
<td>1.45 (0.13)</td>
<td>1.70 (0.10)</td>
</tr>
<tr>
<td>Chl (g m$^{-2}$)</td>
<td>0.27 (0.01)</td>
<td>0.27 (0.02)</td>
<td>0.42 (0.02)</td>
</tr>
</tbody>
</table>

$A_{380}$, net photosynthetic rate at 380 μmol mol$^{-1}$ CO$_2$; $A_{max}$, net photosynthetic rate at 1700 μmol mol$^{-1}$ CO$_2$; $G_s$, stomatal conductance to water vapor; $E$, transpiration rate; $Ls$, stomatal limitation of photosynthesis; $V_{cmax}$, maximum rate of carboxylation; $J_{max}$, maximum rate of electron transport; PNUE, photosynthetic nitrogen use efficiency; LMA, leaf mass per area; $N_{area}$, nitrogen content per unit needle area; Chl, chlorophyll content.

Each value is the mean of eight measurements; the standard error is shown in parentheses.

ANOVA: * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$; n.s. not significant.

For each parameter, values with different letters attached differ significantly, with $P < 0.05$. 

Note: The table includes a brief explanation of each variable, but for clarity, the full text explanation is provided here:

- **$A_{380}$**: Net photosynthetic rate at 380 μmol mol$^{-1}$ CO$_2$.
- **$A_{max}$**: Net photosynthetic rate at 1700 μmol mol$^{-1}$ CO$_2$.
- **$G_s$**: Stomatal conductance to water vapor.
- **$E$**: Transpiration rate.
- **$Ls$**: Stomatal limitation of photosynthesis.
- **$V_{cmax}$**: Maximum rate of carboxylation.
- **$J_{max}$**: Maximum rate of electron transport.
- **PNUE**: Photosynthetic nitrogen use efficiency.
- **LMA**: Leaf mass per area.
- **$N_{area}$**: Nitrogen content per unit needle area.
- **Chl**: Chlorophyll content.