

Seasonal Changes in Root Metabolic Activity and Nitrogen Uptake for Two Cultivars of Creeping Bentgrass

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Abstract. Roots play important roles in plant responses to environmental changes. The objective of this study was to investigate seasonal changes and cultivar variation in root growth, respiratory activity, nitrogen uptake, and carbon allocation in relation to turf performance for two cultivars of creeping bentgrass [*Agrostis stolonifera* L. var. *palustris* (Huds.)] under field conditions. Two cultivars, ‘Penncross’ and ‘L-93’, were managed under USGA-specification putting green conditions, with daily irrigation and mowing at a 3-mm height from May to November in 1999 and 2000. Turf quality of both cultivars declined from the highest rating of 7 to 9 in May to 4 to 5 in August and September, and recovered to above 7 in October and November in both years. This corresponded to seasonal changes in root dry weight, dehydrogenase activity, nitrate reductase (NR) activity, carbon allocation to roots, and ¹⁵N uptake. Compared to Penncross, L-93 generally maintained better turf quality, as well as higher root dry weight, ¹⁵N uptake, NR activity, and carbon allocation during summer months. Previous studies often emphasize the important of a large, extensive root system. The results in the present study demonstrated that root metabolic activities followed the same seasonal pattern and cultivars variation as turf performance, and suggested that decline in root metabolic activities could be contributed to summer decline in turf quality for creeping bentgrass.

The optimum temperature for root growth in cool-season grasses ranges from 10 to 18 °C, and root growth of cool-season grasses decreases as temperature increases above the optimum level (Beard, 1973). Roots of creeping bentgrass exhibited seasonal growth patterns; total root length and number are greatest during the months of March through June, with a second (but smaller) period of increase occurring during October, and roots were smallest during summer (Huang and Liu, 2003; Karnok and Kucharski, 1980; Koski, 1983). Schlossberg et al. (2002) suggested that seasonal cumulative effect of temperature was more influential than instantaneous temperature causing summer root growth decline. Decline in the size of root system during summer for creeping bentgrass may be the result of a greater rate of root dieback than rate of new root production (Huang and Liu, 2003). Small root systems would limit uptake of water and nutrients as well as hormone production.

Most previous studies on seasonal rooting patterns in turfgrass focuses on root growth characteristics (length, weight, or number) (Huang and Liu, 2003; Karnok and Koski, 1983; Kucharski, 1980; Schlossberg et al., 2002; Sweeney et al., 2001), whereas limited information is available on seasonal changes in root metabolic activity and function for cool-season turfgrasses. Previous studies suggested that among root activities, nutrient uptake was one of the most sensitive processes in response

to increasing temperatures in cool-season turfgrass species (Bushoven and Hull, 2001; Huang and Xu, 2000). Nitrogen assimilation largely depends on the activity of nitrate reductase (Bushoven et al., 2002). Nutrient uptake and assimilation were positively related to root respiratory activity and carbon supply from shoots to roots (Greef and Geisler, 1991; Sullivan et al., 2000). Roots are more sensitive to heat stress than shoots, and root growth decline has been shown to proceed shoot growth decline (Xu and Huang, 2000). Disturbances in root metabolic activities under high temperature conditions could lead to deterioration of shoot growth and physiological functions. Therefore, we hypothesized that variation in root metabolic activities are associated with changes in turf performance and differences in heat tolerance in cool-season turfgrasses.

The objective of this study was to investigate seasonal changes and cultivar variation in root respiratory activity, nitrate reductase activity, nitrogen uptake, and carbon allocation for creeping bentgrass. A better understanding of seasonal changes in root activity for creeping bentgrass could help determine timing of implementing management practices to promote active root activities and high quality turf or to alleviate summer decline of root metabolic activity and turf quality.

Materials and Methods

The experiment was conducted on an USGA-specification putting green at the Turfgrass Research Center, Manhattan, Kansas in 1999 and 2000. Two cultivars, ‘L-93’ and ‘Penncross’, were seeded in 205

× 318 cm plots in September 1996. Grass was mowed daily except Sunday at 3 mm in height from early May to early November in 1999 and 2000. During this period, the green was irrigated daily to replace 100% potential evapotranspiration rate, which was measured using mini-lysimeters. The green received four applications (May, July, September, and October) of total nitrogen of 98 kg·ha⁻¹ in 1999 and 65 kg·ha⁻¹ in 2000.

The two cultivars were arranged in a completely randomized block design with three replications. All measurements were taken periodically each month on two subsamples in each plot. Effects of cultivars and time of sampling were determined by analysis of variance according to the general linear model procedure of Statistical Analysis System (SAS Institute, Cary, N.C.). Differences between cultivar means and time of sampling were determined by the least significance difference (LSD) test at the 0.05 probability level.

Daily maximum air temperature (about 150 cm from the ground) was measured using thermocouples connected to a datalogger (CR10X; Campbell Scientific, Inc., Logan, Utah). Data were collected every 10 min and averaged as daily means.

Turf quality was visually rated based on color, uniformity, and density on a 1 to 9 scale, with 9 = the best and 1 = the worst. For the measurement of root dry weight, dehydrogenase activity, and nitrate reductase activity, soil cores of 5 cm in diameter and 20 cm deep were collected from field plots and roots were washed free of soil and collected for analysis. Each parameter was measured for roots collected from two soil cores in each plot.

Root dehydrogenase activity measures respiratory activity, which has been used to differentiate between live and dead tissue using triphenyltetrazolium chloride (TTC) reduction technique (Kniewel, 1973). A 500 mg of fresh roots were placed into test tubes with 0.6% TTC in 0.05 M Na₂HPO₄-NaH₂PO₄ buffer solution. Roots were incubated in a 30 °C water bath in darkness for 20 h, washed twice with water, and then extracted in 95% ethanol at 55 °C for 2 h. Absorbance was measured at 490 nm. Root activity was expressed as absorbance/g fresh weight of roots in 20-mL ethanol solution.

Nitrate reductase activity of roots was determined using the method described by Bushoven and Hull (2001). A 250-mg sample of fresh roots were cut into 5 mm pieces and incubated at 30 °C in 0.1 M phosphate buffer and 1% n-propanol with 0.05 M KNO₃ (pH was 7.5). The production of NO₂ in darkness after 60 min was measured using Grease reagent (1 mL) containing 0.001 g 1-naphtyl-ethylene-diamin, 0.01 g sulphanic acid, and 0.90 g tartaric acid. The amount of NO₂ produced in 60 min was measured spectrophotometrically at 540 nm. Nitrate reductase activity was expressed as nmol NO₂/g/h.

Nitrogen uptake was estimated using the ¹⁵N labeling technique (BassiriRad and Caldwell, 1992). A 22-cm² area of turf was isolated from the surrounding areas (to prevent lateral movement of ¹⁵N enriched NH₄¹⁵NO₃ solution) by installing a plastic tube (5.3 cm in diameter

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and 20 cm deep) in the soil. Fifty ml of 2 mM $\text{NH}_4^{15}\text{NO}_3$ solution (5% of N atom enrichment) was slowly injected into the soil at 3 and 6-cm depths in the isolated soil column. Four days after labeling, the labeled plants were harvested along with 20-cm deep soil core. Roots were washed free of soil and were killed at 105 °C and dried at 70 °C. Total N and ^{15}N content in roots was determined. The ^{15}N content was assumed to be a measure of root N uptake within the labeling (4 d) period (Thornton and Millard, 1996). Specific ^{15}N content ($\text{ng}\cdot\text{g}^{-1}$ dry weight) in roots was calculated using the following equation described in Millard and Neilsen (1989): $(\text{atom}\% \text{ of sample } ^{15}\text{N} - 0.367) \times 10000 / \text{dry weight}$. The atom% of sample ^{15}N was measured using a mass spectrophotometer (Shimadzu Scientific Instruments Inc. Columbia, S.C.).

Carbon allocation to roots was determined using pulse $^{14}\text{CO}_2$ labeling technique as described by Huang and Fu (2000). Plant canopy was enclosed in a clear plexiglass chamber (5 cm tall and 10 cm in diameter) and exposed to 10 $\mu\text{Ci } ^{14}\text{CO}_2$ for 20 min. Three days after $^{14}\text{CO}_2$ labeling, shoots and roots were washed free of soil, killed at 105 °C for 30 min, and dried at 75 °C for 48 h. Twenty milligrams of sample was digested with 4 mL solubilizing liquid (Solvable; Dupond, Wilmington, Del.) at 50 °C for 24 h and bleached with 0.1 mL of 30% H_2O_2 for 1 h at 25 °C (Huang et al. 1993). Fifteen milliliters of scintillation cocktail

for ^{14}C counting were added to the solutions and were measured in a scintillation counter (Packard, Deers Grove, Ill.). The proportion of newly photosynthesized carbon that was allocated to roots was calculated.

Results and Discussion

Seasonal changes in air temperatures. Daily maximum air temperature exhibited clear seasonal patterns in both 1999 and 2000 (Fig. 1). It was averaged 24 °C in May and increased to the highest level in July 1999 and August 2000. Highest temperature was 40 °C at the end of July in 1999 and 43 °C at the end of August 2000. By October and November in both years, air temperature declined to below 24 °C, similar to the level in May.

Seasonal changes in turf quality, root growth, and metabolic activities. Turf quality for L-93 and Penncross were at the highest level of 8.3 and 7.6 in May and June, respectively; the rating then declined to the lowest level of 5.8 and 5 in August 1999 (Fig. 2A) and of 6.0 and 4.0 in September 2000 for L-93 and Penncross, respectively (Fig. 2B). Turf quality recovered to an average of 8.6 and 7.1 in October and November in both years for L-93 and Penncross, respectively. Root dry weight also exhibited seasonal changes for both cultivars in both years (Fig. 3A and B), with a cultivar average of 40% decline

from May to August in 1999 and 63% from May to September 2000. The changes in root dry weight did not follow the same pattern as for turf quality in 1999. Previous studies have shown that turf quality decline during summer for creeping bentgrass was accompanied by a decline in root growth characteristics, and the timing of minimum root length, biomass, and number corresponded to the onset of turf quality decline in creeping bentgrass (Huang and Liu, 2003; Schlossberg et al., 2002; Sweeney et al., 2001). Our study also found that summer turf quality decline of creeping bentgrass may not only be related to decreases in root size as biomass or length, but also could be associated with the deterioration of root metabolic activities in terms of carbon supply or demand, dehydrogenase activity, and nitrogen uptake and assimilation, as discussed in the following sections. Seasonal changes in turf quality and root growth were closely related to seasonal temperature changes (Fig. 1), although other environmental factors such as relative humidity may also affect these parameters. Lowest turf quality (Fig. 2), root growth and activities (Fig. 3–7) were observed when air temperature reached the maximum in July and August in both years (Fig. 1), suggesting that temperature had significant impact on turf and root growth.

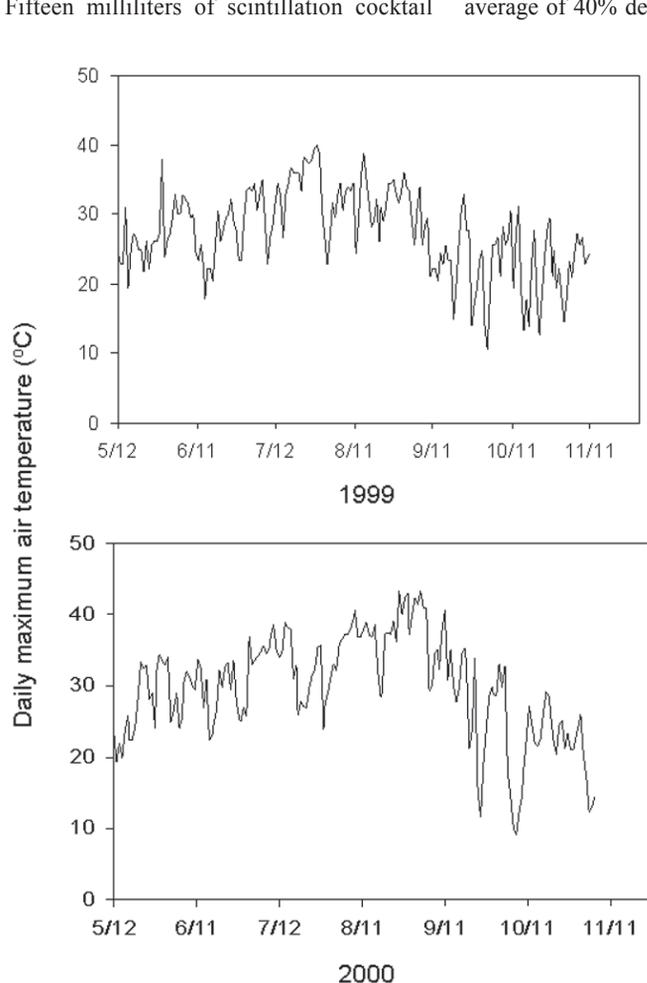


Fig. 1. Daily maximum air temperature from May to November in 1999 and 2000.

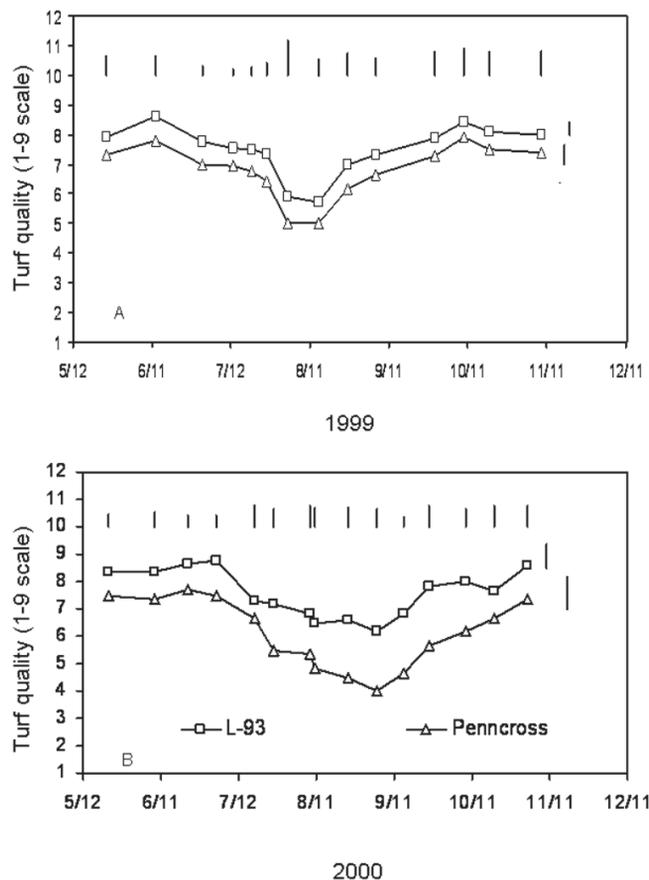


Fig. 2. Seasonal changes in turf quality for two creeping bentgrass cultivars, L-93 and Penncross, in 1999 (A) and 2000 (B). Turf quality was visually rate on the scale of 1 to 9, with 9 = the best. Vertical bars on the top indicate LSD values ($P < 0.05$) for cultivar comparison at a given date of year. Vertical bars on the right indicate LSD values ($P < 0.05$) for seasonal changes for a given cultivar.

¹⁴Carbon allocation to roots for both cultivars was lowest in July and August 1999, accounting for 10.4% of total carbon fixation; the proportion of ¹⁴Carbon in roots was highest (15%) in May 1999 and October (13%) (Fig. 4A). Similar patterns were found in 2000, when the proportion of ¹⁴Carbon in roots was 23%, 11%, and 20% in June, September, and November 2000, respectively (Fig. 4B). The summer decline in ¹⁴C allocation to roots was more pronounced for Penncross than for L-93 in both years. The decrease in the proportion of ¹⁴C allocation to roots suggested that the amount of newly photosynthesized carbon allocated to roots decreased in the summer, which could be related to the decline in root dry weight in summer months. This result also indicated that root demand for carbon or carbon metabolic activity decreased during summer months, because carbon supply from shoots to roots is largely dependent on the demand or activity of a root system (Hopkins, 1999).

Dehydrogenases are critical mitochondrial enzymes involved in energy production in respiration that are used in growth, nutrient uptake, and tissue maintenance (Hopkins, 1999). Dehydrogenase enzyme activity is a general measure of tissue respiratory activity, and has been used to quantify root respiratory activity or viability (Knievel, 1973). Some dehydrogenases such as glutamate dehydrogenase are also involved in nitrogen metabolism (Beevers and Hageman, 1969). In the present

study, root dehydrogenase activity for both cultivars decreased to the lowest level in August 1999 (only 33% of the level in May) and then increased in October through November to 85% of the level in May (Fig. 5A). In 2000, dehydrogenase activity decreased to the lowest level in late June, remained low through September, and then recovered in October (Fig. 5B). These results indicated that energy production through respiration and nitrogen metabolism in roots for creeping bentgrass were limited during summer months. Nelson and Treharne (1973) reported that activity of the respiratory enzymes such as malate dehydrogenase and isocitrate dehydrogenase in tall fescue (*Festuca arundinacea* Schreb.) decreased with increasing temperatures. Decreased dehydrogenase activity has been associated with heat stress damages (Kaur and Nainawatee, 1989).

Nitrogen uptake and metabolism is the main limiting step in amino acid and protein production, which has significant impact on plant response to environmental stresses (Beevers and Hageman, 1969). In 1999, ¹⁵N content was evaluated in August and October (Table 1). ¹⁵N content was 72% higher in October 1999 than that in August for both cultivars. In 2000, ¹⁵N content was measured monthly from May to October (Fig. 6). The content of ¹⁵N was highest in May, decreased to the lowest level in July and August, and increased in October for both cultivars. Nitrate reductase is the key enzyme

in nitrate assimilation and plant development (Beevers and Hageman, 1969), and is sensitive to high temperatures (Lauri and Stewart, 1993; Onwueme et al., 1971). Nitrate reductase activity of roots was measured monthly from May to October in 2000 (Fig. 7). Lowest NR activity was observed in August for L-93 and in September for Penncross. Both ¹⁵Nitrogen content and NR activity decreased during summer. These data indicated that both root nutrient uptake and assimilation activities were inhibited when creeping bentgrass experienced high temperatures during summer. Younis et al. (1965) found that an increase in temperature from 30 to 35 °C caused a 60% to 70% decrease in NR activity in young corn (*Zea mays* L.) plants. The reduction in nitrogen uptake and assimilation activities under high temperatures could be related to the limited carbon supply to roots during summer as described above. Sullivan et al. (2000) demonstrated that the ability of roots to assimilate nitrate depends on carbohydrate supply from shoots in tall fescue. Greef and Geisler (1991) also found a high level of sugars induced a high nitrogen uptake in corn.

Cultivar variation in turf quality, root growth, and metabolic activities. Previous studies demonstrated that L-93 was more heat tolerant than Penncross (Xu and Huang 2000, 2001). Similar to previous reports, turf quality was higher for L-93 than for Penncross on most measurement dates in both years, rang-

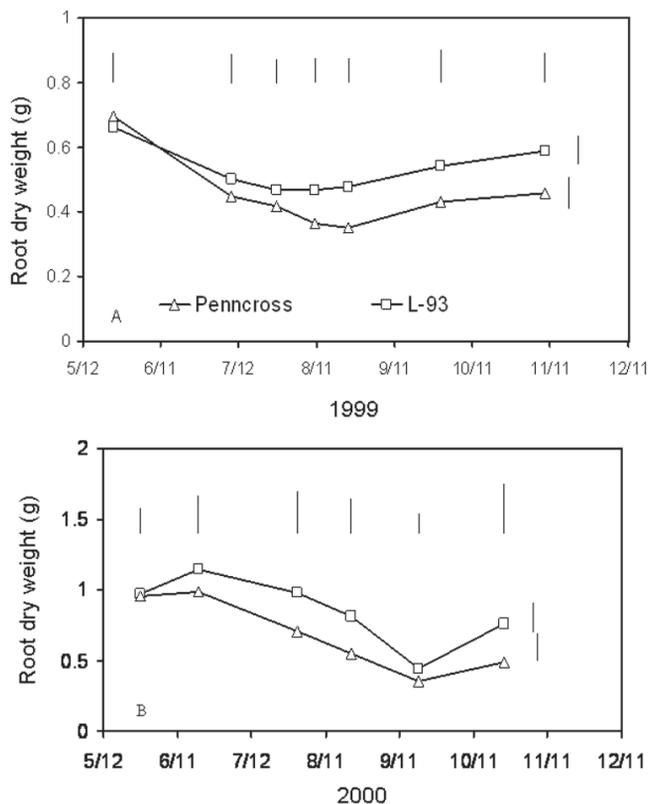


Fig. 3. Seasonal changes in root dry weight for two creeping bentgrass cultivars, L-93 and Penncross, in 1999 (A) and 2000 (B). Root dry weight was the weight of roots collected from a 393 cm³ soil in the surface 20-cm soil. Vertical bars on the top indicate LSD values ($P < 0.05$) for cultivar comparison at a given date of year. Vertical bars on the right indicate LSD values ($P < 0.05$) for seasonal changes for a given cultivar.

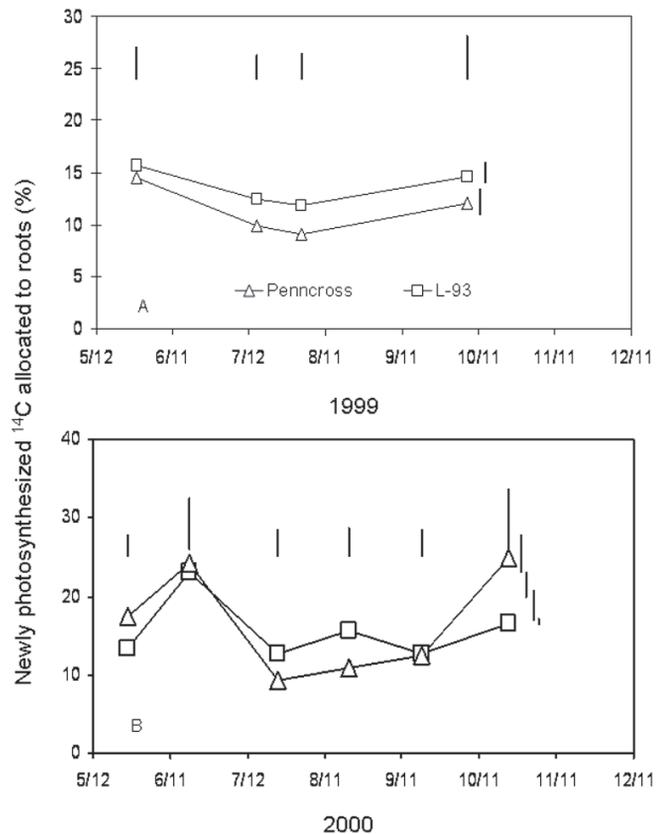


Fig. 4. Seasonal changes in carbon allocation to roots for two creeping bentgrass cultivars, L-93 and Penncross, in 1999 (A) and 2000 (B). Vertical bars on the top indicate LSD values ($P < 0.05$) for cultivar comparison at a given date of year. Vertical bars on the right indicate LSD values ($P < 0.05$) for seasonal changes for a given cultivar.

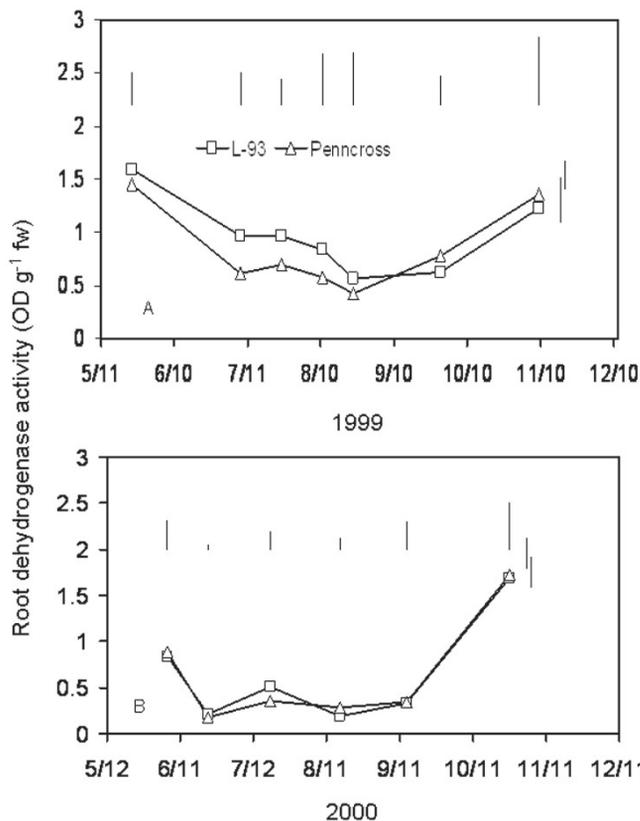


Fig. 5. Seasonal changes in root dehydrogenase activity for two creeping bentgrass cultivars, L-93 and Penncross, in 1999 (A) and 2000 (B). Vertical bars on the top indicate LSD values ($P < 0.05$) for cultivar comparison at a given date of year. Vertical bars on the right indicate LSD values ($P < 0.05$) for seasonal changes for a given cultivar.

ing from 5.7 to 8.6 for L-93 and from 5.0 to 7.8 for Penncross in 1999 and from 6.2 to 8.6 for L-93 and 4.0 to 7.5 for Penncross in 2000 (Fig. 2A and B). Cultivar difference was most pronounced during summer months in 2000 (Fig. 2B). L-93 had a greater root dry weight than Penncross from August to November 1999 and in July and August 2000. Lehman and Engelke (1991) reported that creeping bentgrass cultivars that produce more roots and deeper roots were better able to survive intensive heat. Bonos and Murphy (1999) found that heat-tolerant Kentucky bluegrass (*Poa pratensis* L.) cultivars had more roots than heat-sensitive cultivars.

Development of deep, extensive root systems (larger root mass or length) is often emphasized as an important factor in plant growth and adaptation to environmental stresses. Cultivar differences in turf quality dur-

ing summer could also be associated with greater root metabolic activities. The proportion of newly photosynthesized carbon allocated to roots was 29% higher in L-93 than in Penncross in July and August in both years (Fig. 4A, B), reflecting higher carbon metabolism in roots of L-93. L-93 also had higher dehydrogenase activity than Penncross in July 1999 (Fig. 5A); however, no significant difference in root dehydrogenase activity was detected between the two cultivars in 2000 (Fig. 5B). Root ¹⁵N content was 29% and 21% higher in L-93 than in Penncross in August and October 1999, respectively (Table 1); in 2000, ¹⁵N content was also higher for L-93 than for Penncross in August and September (Fig. 6). L-93 also had a greater NR activity than Penncross in September 2000 (Fig. 7). These results suggested roots of L-93 not only were able to maintain higher nitrogen uptake and assimilation activities than roots of Penncross during summer months. The larger root system in combination with high metabolic activities could contribute to better heat tolerance for creeping bentgrass.

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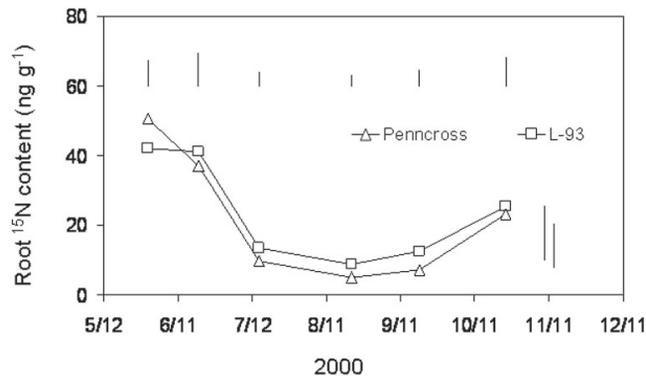


Fig. 6. Seasonal changes in ¹⁵N uptake for two creeping bentgrass cultivars, L-93 and Penncross, in 2000. Data for 1999 was presented in Table 1. Vertical bars on the top indicate LSD values ($P < 0.05$) for cultivar comparison at a given date of year. Vertical bars on the right indicate LSD values ($P < 0.05$) for seasonal changes for a given cultivar.

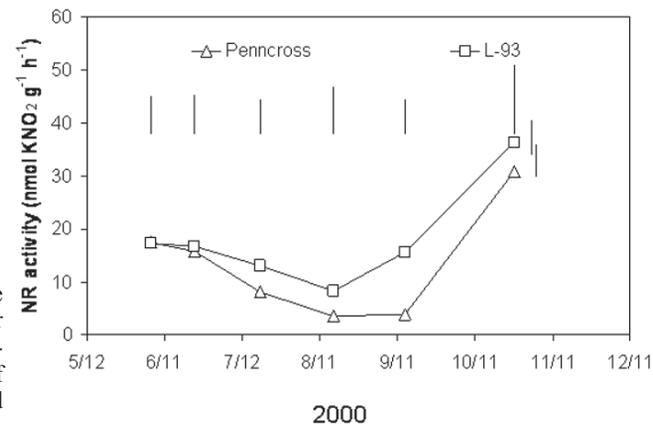


Fig. 7. Seasonal changes in nitrate (NR) reductase activity for two creeping bentgrass cultivars, L-93 and Penncross, in 2000. Vertical bars on the top indicate LSD values ($P < 0.05$) for cultivar comparison at a given date of year. Vertical bars on the right indicate LSD values ($P < 0.05$) for seasonal changes for a given cultivar.

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Table 1. Root ¹⁵N uptake for two cultivars of creeping bentgrass, L-93 and Penncross, in August and October 1999. Data for 2000 was presented in Fig. 6.

Cultivar	¹⁵ N (ng·g ⁻¹ dry wt)	
	3 Aug.	8 Oct.
L-93	44.02 aB ²	73.53 aA
Penncross	34.01 bB	60.61 bA

²Means followed by the same lower case letters within a column or the same upper case letters within a row were not significantly different based on LSD test at $p = 0.05$.

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