Gypsum Effects on Growth and Macroelement Uptake of Field-grown Asimina triloba (Pawpaw) Irrigated with Low-saline, Sodic Water

G.A. Picchioni¹
Department of Agronomy and Horticulture, New Mexico State University, Las Cruces, NM 88003

C.J. Graham
Louisiana State University Agricultural Center, Calhoun Research Station, Box 539, Calhoun, LA 71225

A.L. Ulery
Department of Agronomy and Horticulture, New Mexico State University, Las Cruces, NM 88003

Abstract. Asimina triloba (L.) Dunal is an underused tree species with demonstrated potential as a new fruit crop and landscape ornamental plant. Best management practices for A. triloba are not adequately defined, particularly for field establishment in high-Na conditions characteristic of numerous southern U.S. production areas. We evaluated the growth and net macrolelement uptake of field-grown A. triloba seedlings on soil amended with a single addition of gypsum at 0, 7.5, or 15 t ha⁻¹ and later receiving a regular supply of Na-affected but nonsaline irrigation water [sodium adsorption ratio (SAR) of 15.5 and electrical conductivity (EC) at 0.4 dS·m⁻¹]. Over two growing seasons, the soil saturation extract Ca concentration increased while the soil saturation extract SAR decreased with increasing gypsum rate. Amending the soil with gypsum increased total lateral branch extension per tree by 60% to 73% and trunk cross-sectional area (TCSA) per tree by 68% to 87% above a non-gypsum-amended control treatment. Total dry matter accumulation and the net uptake of N, P, and K per tree were over 100% greater following gypsum application as compared to controls. The growth and mineral uptake-enhancing effects of gypsum were likely related to functions of Ca at the root level and on soil physical properties that should be considered in establishing young A. triloba trees with irrigation water containing high sodicity but relatively low total salinity.

Asimina triloba (pawpaw) has received considerable attention as a new commercial tree fruit crop, landscape ornamental, and medicinal plant (Callaway, 1992; Huang et al., 2003; Layne, 1996; Zhao et al., 1993). However, difficulty in transplanting A. triloba is a barrier toward successful commercialization of the species (Callaway, 1990; Callaway and Callaway, 1992; Darrow, 1975). Several short-term pot culture fertilization studies have been conducted on A. triloba (Pomper et al., 2002a, 2000b, and 2000c), but there are no quantitative data on factors affecting its growth and mineral uptake characteristics in field conditions. This lack of data is significant, since young A. triloba trees are reported to be inherently slow-growing with slowly developing root systems (Darrow, 1975). Enhancement of root growth would speed nursery or orchard establishment, grafting and budding of fruit-bearing cultivars, and regeneration of renewal biomass for extraction of natural products.

High-Na (sodicity) is a transcontinental concern in the southern latitudes of the United States, where it threatens a diversity of existing commercialized tree fruit crops, such as pecan, pistachio, blueberry, citrus, plum, and peach (Karakas et al., 2000; Picchioni et al., 1990, 2000; Wright et al., 1992; Zekri and Parsons, 1990; Ziska et al., 1991). Northern Louisiana is on the southwestern fringe of the A. triloba indigenous zone, where groundwater sodium adsorption ratio (SAR) may exceed 40 and EC, 9 dS·m⁻¹ (U.S. Geological Survey, 1991). Numerous groundwater monitoring stations in northern Louisiana report high salinity combined with ionic imbalances, most notably cases in which Na dominates the cationic fraction, while Mg and Ca concentrations are at trace levels to below 1 meq L⁻¹ (U.S. Geological Survey, 1991). Local groundwater supply during the present study in Ruston (described below) had an average SAR of 15.5 but relatively low average total salinity, expressed as EC, at 0.4 dS·m⁻¹.

Reducing potential sources of environmental stress is essential for successful commercialization of new crops (Wallis et al., 1989). In reference to the present study, the inherent difficulty in field establishment of A. triloba could be exacerbated with Na-affected irrigation waters and soils. Injurious effects of the Na-dominated irrigation waters of northern Louisiana were highlighted in a recent study involving young trees of Craepean opacu Hook. and Arn. (Picchioni and Graham, 2001). Like A. triloba, C. opacu is indigenous to the North American temperate zone and under consideration as a new fruit crop and ornamental tree.

Calcium serves an important role in countering negative effects of Na on irrigation waters, soils, and plants. In addition to its functions at the root level (Epstein, 1961; Hanson, 1984; Picchioni et al., 1991), Ca maintains stable, flocculated clay mineral aggregates that are essential for good soil structure and internal drainage (Bohn et al., 1985). Gypsum is the most widely used neutral Ca salt for maintaining or amending soil structure in Na-affected soils (Mengel and Kirkby, 1987). We hypothesized that gypsum would improve field establishment of A. triloba irrigated with sodic, nonsaline water. Therefore, our objective was to evaluate gypsum effects on A. triloba field establishment and the net uptake of macrolelements under irrigated conditions of high sodicity and relatively low total salinity.

Materials and Methods

Preparation of seedling tree material. Seeds of A. triloba were collected in Fall 1993 from fruit of trees growing in the wild along the Mississippi River floodplain and just east of the town of Transylvania in northeastern Louisiana. Seeds were separated from the fruit, rinsed with water, and subjected to cold-moist stratification as prescribed by Layne (1996). After stratification, seeds were sown individually at a depth of 2 cm in 17-L plastic pots (27 cm wide x 30 cm tall) filled with pine bark. Seedlings were grown outdoors at the Calhoun Research Center under 50% shade and irrigated with tap water (EC of 0.4 to 0.5 dS·m⁻¹) as needed for 16 months until Spring 1995 field planting (described below). Using a photometer (General Electric Model 214), the summer maximum photosynthetic photon flux (PPF) under the shaded conditions was estimated to be 750 to 780 μmol·m⁻²·s⁻¹. Photometric units were converted to quantum units by the method described by Thimijan and Heins (1983). There were 179 cm of rainfall over the 16 months of shaded pot culture and the air temperature ranged from –9 to 36 °C (lowest winter minimum and highest summer maximum temperatures, respectively), according to an on-site weather station.

Fifty-four dormant, unbranched, 16-month-
Field site description and cultivation. A 0.05-ha field plot at the Louisiana Tech Univ. horticultural farm in Ruston (≈17 km west of the Calhoun Research Center) was used for the study. The site was fallow before use in the present investigation. The soil was a Ruston fine loamy, siliceous, semiactive, thermic, Typic, Paleudult, with a brown fine sandy loam in the surface and subsurface layers (to ≈15 cm), underlain by red clay loam in the upper subsoil. The middle and lower subsoil were comprised of yellowish fine sandy loam and red sandy clay loam, respectively. During the field study growing seasons of 1995 and 1996, the air temperature ranged from –13 to 41 °C (lowest winter minimum and highest summer maximum temperatures, respectively). The summer maximum PFD (estimated as described above) was 1450 to 1500 µmol·m−2·s−1.

Before planting, raised beds measuring 0.38 m in height and 0.9 m in top width were prepared from the top 15 cm of fine sandy loam scraped from the 4.6-m middle aisles between the matted beds (beds). Soil organic matter content was determined by the methods described by Nelson and Sommers (1996), averaged 1.0% in this portion of the profile, and did not change throughout the study. Beds were initially shaped and packed by making three passes with a 23-cm-diameter roller, then gypsum (CaSO4·2H2O; pelletized agricultural grade) was added to the bed tops to a depth of 15 cm to provide CaSO4 at 0.0 (control), 7.5, or 15.0 t·ha−1. The single application of gypsum was made on 3 Apr. 1995, and the rates selected from within the range recommended for agricultural crops (California Fertilizer Association, 1995; Mengel and Kirkby, 1987). Following gypsum application, beds were again packed with the roller. The three application rates were arranged in a randomized complete-block design with three trees per replication and six replications, totaling 18 trees per treatment. The trees were then transplanted on 7 Apr. 1995 atop the beds at a spacing of 5.5 m between bed centers and 1.5 m within the bed row (1212 trees per ha), and watered thoroughly. Each three-tree replication measured 4.5 m long × 0.9 m wide. Trees were grown up to their excavation on 17 Oct. 1996 and had experienced two growing seasons in the field in addition to a single growing season (16 months) of pot culture.

Throughout the duration of the study, irrigation was supplied through a manually-operated drip irrigation system with two emitters per tree, each with a flow rate of 4.5 L·h−1. Emitters were positioned on top of the bed center and at a lateral distance of 30 cm from the trunks. Each irrigation supplied ≈53 L water per tree, which was estimated to be sufficient to fill the entire pore volume of the fine sandy loam bed. The irrigation source (tap water) had an EC of 0.4 dS·m−1, SAR of 15.5, pH of 7.7, and ion concentrations (in meq·L−1) as follows: Na (4.7), Ca (0.14), Mg (0.04), Cl (1.0), HCO3− + CO3− (3.0), and SO4− (16.1). No field irrigation guidelines have been established for A. triloba. Therefore, irrigation planning was aided by general orchard irrigation methods as prescribed for similar growing conditions of east Texas by Worthington et al. (1986), while considering local monthly grand average U.S. Weather Service peak pan evaporation data over a prior 20-year period. Local monthly average pan evaporation values for the 1995 and 1996 growing seasons were within 10% of their previous 20-year averages. The combined natural precipitation and irrigation met 84% of the 1995 growing season and 70% of the 1996 growing season as compared to the same months of 1995 as compared to the same months of 1995 (76 vs. 42 cm, respectively).

On three dates during the study (5 May and 24 June 1995, and 28 Apr. 1996), Peters 20N–4.4P–16.6K water-soluble fertilizer (J.R. Peters, Allentown, Pa.) was added to the irrigation solution using a fertilizer injector (Gewau model 6, Hermann A. Wirth, Princeton, Fla.). Each fertigation solution contained (in mg·L−1) 146–32–121 K, and delivered N, P, K at 50, 11, and 42 kg·ha−1, respectively. On the final fertigation, Mg was supplied (as MgSO4·7H2O) to all treatments at 50 mg·L−1 to provide an apparent Mg deficiency. A high concentration of basal shoot leaves induced by the 7.5 and 15.0 t·ha−1 gypsum treatments and noted at the end of the 1995 growing season. The EC of each fertigation solution averaged 1.5 dS·m−1 and pH ranged from 6.4 to 6.7.

Field sampling, data collection, and analysis. Soil samples were collected on 2 Apr. 1995 (1 d before gypsum application), 28 Apr. 1996 (following the first growing season), and 16 Oct. 1996 (following the second growing season). Two soil cores measuring 20 cm in depth and 2.5 cm in diameter were taken from the midpoints of the outer and middle trees of each replication. Soils were dried at 60 °C for 2 weeks and passed through a 1-mm sieve. A 100-g subsample was used to prepare the soil saturation extract (U.S. Salinity Laboratory Staff, 1954), and the pH and EC were determined, followed by the concentrations of Na, Ca, and Mg using an inductively-coupled plasma emission (ICP) spectrometer (JY70+, Instruments S.A., Edison, N.J.). Total Kjeldahl N (TKN) concentration was determined using an autoanalyzer (model II, Technicon Instru-
increases in tree height, lateral branch length, TCSA, biomass, and macromolecule content, and for the tree visual rating, the average value of three trees comprised a replication. The ground tissues from the three trees per replication were pooled for mineral analysis; thus, mineral-related data are the means of six analytical determinations per treatment. Macromolecule content per organ at the time of tree excavation (17 Oct. 1996) was determined as the product of the three-tree average organ dry weight and the three-tree average macromolecule concentration of the excavated trees. Average macromolecule content per organ at planting was also determined (as above), and then subtracted from the value obtained at excavation to determine net macromolecule uptake per tree during the time of gypsum treatment, discounting the net uptake by leaves. Where significant, sums of squares for gypsum application main effects were partitioned into single degree of freedom linear and quadratic orthogonal contrasts.

Results

Soil compositional changes. One day before gypsum application, soil saturation extract pH, EC, and cationic characteristics did not differ between the marked treatment experimental units (2 Apr. 1995 sampling date in Table 1). At this time, TKN concentration ranged between 550 and 589 mg·kg–1 dry weight, P between 39 and 40 mg·kg–1 dry weight, and K between 20 and 26 mg·kg–1 dry weight, with no differences between the treatment plots (data not shown). Thereafter, soil saturation extract pH, and the concentrations of TKN, P, and K were unaffected by gypsum rate throughout the study period (data not shown).

The soil saturation extract EC (ECe) of the control treatment (gypsum at 0.0 t·ha–1) was identical at each sampling date, but at the second sampling date (13 months after gypsum application), ECe had increased by 1.8 to 3.4 times because of gypsum application (Table 1). By the end of the study (18 months after transplanting (Table 1). This trend had already become evident near the close of the first growing season (average TCSA increases of 49, 63, and 71 mm2/tree by 29 Sept. 1995, for 0.0, 7.5, and 15.0 t·ha–1 gypsum rates, respectively). In addition to the linear increases in lateral branch extension and TCSA with gypsum rate, the level of tree health and vigor (as visually assessed by numerical ratings) also improved with gypsum rate (visual rating value became smaller with increasing gypsum rate) at the end of both 1995 and 1996 growing seasons (Table 2).

The increases in TCSA over both growing seasons combined were 68% to 87% greater if gypsum was previously added to the soil as compared with no gypsum application (Table 2). This trend had already become evident near the close of the first growing season (average TCSA increases of 49, 63, and 71 mm2/tree by 29 Sept. 1995, for 0.0, 7.5, and 15.0 t·ha–1 gypsum rates, respectively). In addition to the linear increases in lateral branch extension and TCSA with gypsum rate, the level of tree health and vigor (as visually assessed by numerical ratings) also improved with gypsum rate (visual rating value became smaller with increasing gypsum rate) at the end of both 1995 and 1996 growing seasons (Table 2).

Table 1. Soil saturation extract pH, electrical conductivity (EC), selected cations, and sodium adsorption ratio (SAR) to a depth of 20 cm before and during 18 months of gypsum treatment and field establishment of A. triloba trees.  

<table>
<thead>
<tr>
<th>Sampling date</th>
<th>Gypsum rate (t·ha–1)</th>
<th>Soil saturation extract characteristics</th>
<th>meq·L–1</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>pH</td>
<td>EC (dS·m–1)</td>
<td>Mg</td>
</tr>
<tr>
<td>2 Apr. 1995</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0.0</td>
<td>5.24</td>
<td>0.55</td>
<td>0.77</td>
</tr>
<tr>
<td>7.5</td>
<td>5.15</td>
<td>0.52</td>
<td>0.71</td>
</tr>
<tr>
<td>15.0</td>
<td>5.08</td>
<td>0.48</td>
<td>0.60</td>
</tr>
<tr>
<td>Significance</td>
<td>NS</td>
<td>NS</td>
<td>NS</td>
</tr>
<tr>
<td>28 Apr. 1996</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0.0</td>
<td>5.08</td>
<td>0.55</td>
<td>0.25</td>
</tr>
<tr>
<td>7.5</td>
<td>5.26</td>
<td>0.97</td>
<td>0.95</td>
</tr>
<tr>
<td>15.0</td>
<td>5.21</td>
<td>1.87</td>
<td>2.48</td>
</tr>
<tr>
<td>Significance</td>
<td>NS</td>
<td>*</td>
<td>**</td>
</tr>
<tr>
<td>16 Oct. 1996</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0.0</td>
<td>5.31</td>
<td>0.55</td>
<td>0.07</td>
</tr>
<tr>
<td>7.5</td>
<td>5.41</td>
<td>0.55</td>
<td>0.11</td>
</tr>
<tr>
<td>15.0</td>
<td>5.34</td>
<td>0.90</td>
<td>0.82</td>
</tr>
<tr>
<td>Significance</td>
<td>NS</td>
<td>*</td>
<td>NS</td>
</tr>
</tbody>
</table>

Each value is the mean of six replications.

*Gypsum was applied on 3 Apr. 1995, trees were planted on 7 Apr. 1995, and excavated for analysis on 17 Oct. 1996.

*Calculated as Na/(Ca + Mg/2)1/2, all ions in meq·L–1.

NS, *, **Non-significant or significant at P ≤ 0.05 or 0.01, respectively; linear (L) or quadratic (Q) orthogonal contrasts partitioned from the sums of squares for main effect of gypsum rate.

Table 2. The effects of gypsum application on 3 Apr. 1995 on increases in A. triloba tree height, total lateral branch extension, and trunk cross-sectional area (TCSA), from 7 Apr. 1995 to 17 Oct. 1996. Trees were visually rated in Fall 1995 and 1996.

<table>
<thead>
<tr>
<th>Gypsum rate (t·ha–1)</th>
<th>Increase in tree height (cm)</th>
<th>Lateral branch extension (cm)</th>
<th>Increase in TCSA (mm2)</th>
<th>Visual tree rating</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.0</td>
<td>38.7</td>
<td>63.8</td>
<td>99.0</td>
<td>3.8</td>
</tr>
<tr>
<td>7.5</td>
<td>49.3</td>
<td>102.2</td>
<td>185.2</td>
<td>2.8</td>
</tr>
<tr>
<td>15.0</td>
<td>49.8</td>
<td>110.4</td>
<td>166.5</td>
<td>2.5</td>
</tr>
<tr>
<td>Significance</td>
<td>NS</td>
<td>NS</td>
<td>NS</td>
<td>NS</td>
</tr>
</tbody>
</table>

Each value is the mean of six replications.


NS, *, **Non-significant or significant at P ≤ 0.05 or 0.01, respectively; linear (L) or quadratic (Q) orthogonal contrasts partitioned from the sums of squares for main effect of gypsum rate.
Depending on gypsum rate, total dry weight per tree at the time of tree excavation had increased by 15- to 33-fold above the average value at transplanting. Final dry weights were variable within treatments. The average CV values across the gypsum rates for dry weight of roots, trunk plus lateral branches, and their combined totals were 30.9%, 44.4%, and 32.6%, respectively. Trunk plus lateral branches comprised a relatively small fraction of the total excavated tree dry weight (23% across the treatments).

During the 18 months after gypsum application, over twice the amount of dry matter had accumulated in tops and roots of trees growing on gypsum-amended soil as compared to trees on the non-amended soil (Fig. 1A). Total dry matter gain per tree was 79 to 84 g higher under the gypsum-amended conditions in relation to controls. From 71% to 78% of the difference in total tree dry matter gain between trees on gypsum-amended soil and those not on gypsum-amended soil was attributed to the difference in root dry matter accumulation. Linear contrasts for dry matter accumulation in trunk plus lateral branches, roots, and their combined totals were significant at \( P \leq 0.05 \), even though there was little variation in organ dry matter gains between the 7.5 and 15.0 t·ha\(^{-1} \) gypsum treatments.

**Macroelement concentrations and net uptake per tree.** At the time of tree excavation, none of the macroelement concentrations (N, P, K, Ca, Mg, and S expressed as percentage of dry weight) in roots and combined trunk plus lateral branches were affected by gypsum rate (Table 3). The average CV (across the treatments) ranged from 7% to 18%.

Because of the gypsum rate effects on tree biomass accumulation but similarity in mineral concentrations between the gypsum rates, the net uptake of N, P, and K by roots and combined trunks plus lateral branches followed the same trend with gypsum rate as did the accumulation of dry matter in these tissues (Fig. 1B–D). Identical patterns were observed for net Ca, Mg, and S uptake (data not shown). High variability in total net N, P, and K uptake per tree within the gypsum rates (CV average of 32% to 39%, depending on the element) resulted primarily from the relatively high variability in biomass accumulation, and to a lesser degree from variability in macroelement concentrations within gypsum rates, as noted above.

Following transplanting, the net uptake of N, P, and K per tree (trunk, lateral branches, and roots) was equivalent to levels that were 10- to 21-fold (N), 11- to 24-fold (P), and 7- to 15-fold (K) those measured in the trees at the time of transplanting. Gypsum rate did not significantly affect the net uptake of N, P, and K by trunks plus lateral branches (Fig. 1B–D), nor did it affect total net macroelement uptake (roots + trunks + lateral branches) when expressed as a function of the unit increase in root dry weight between transplanting to excavation (data not presented). However, total net N, P, and K uptake per tree increased with gypsum rate, because of the large influence of gypsum on the accumulation of biomass.

**Fig. 1.** Influence of gypsum rate on net dry matter (DM) accumulation (A) and the net uptake of N, P, and K (B, C, and D, respectively) by trunks, lateral branches, and roots during 18 months of field establishment of *A. triloba*. Significance (*) or nonsignificance (NS) of linear orthogonal contrast partitioned from the main effects of gypsum rate (\( P \leq 0.05 \)). Each value is the mean ± se of six replications. Initial values at transplanting: root dry weight (3.7 g per tree); trunk plus lateral branch dry weight (0.9 g per tree); root N, P, and K content (66.0, 9.3, and 59.7 mg per tree, respectively); trunk plus lateral branch N, P, and K content (15.7, 1.5, and 5.6 mg per tree, respectively).
During the course of the final growing season, there was a sharp reduction in soil saturation extract Ca concentration by increasing the soil saturation extract Ca concentration by as much as 13 times, depending on gypsum rate and time following the application. This long-established effect of gypsum is known to increase stability (e.g., flocculation) of clay mineral aggregates and thereby improve soil permeability, aeration, and internal drainage in Na-affected soils (Bohn et al., 1985).

During the course of the final growing season, there was a sharp reduction in soil saturation extract Ca concentration with a marginal increase in soil saturation extract SAR with the highest gypsum rate (15.01 t·ha⁻¹). This indicates that repeated gypsum applications may be necessary beyond the conditions of our study to maintain a favorable soil condition for A. triloba orchard management with Na-affected irrigation water. The apparent Mg deficiency observed after the first growing season (gypsum-treated trees only) may have resulted from Ca antagonism of Mg uptake (Mengel and Kirkby, 1987). Accordingly, in irrigated conditions combined with gypsum application on soils of low-Mg availability (as in our study), Mg fertilization may be essential for this species. Coincidentally, the higher soil saturation extract Mg concentrations of the gypsum-amended soil at the second sampling date had a negligible effect on soil saturation extract SAR, and may have resulted from impurities in the agricultural grade gypsum.

Asimina triloba is reported to possess a dominant taproot that may be adaptive for exploring the soil profile in its indigenous riparian habitats (Nash and Graves, 1993). The degree to which soil can be penetrated by roots is dependent on the soil-related physical functions of Ca listed above (Mengel and Kirkby, 1987; Simpson et al., 1997), and in the present study, variation in A. triloba root growth comprised over 70% of the difference in tree biomass accumulation between trees on amended and non-amended soil. Root growth is also directly dependent on Ca in the soil solution (Hanson, 1984; Piccioni et al., 2001; Pooviah and Reddy, 1991). While the Ca requirement of A. triloba is not adequately known, the saturation extract Ca concentration at 15.0 t·ha⁻¹ gypsum was in the range of a calcareous soil (Piccioni et al., 1990). We did not determine the extent of Ca transport or root penetration into the dominant red clay subsoil, where gypsum would be expected to exert a significant influence on the physical properties of Na-affected soil. Further study is needed to evaluate variation in A. triloba root growth restriction in Na-affected, high-clay soils of varying Ca status.

Numerous A. triloba cultivars have originated from Ohio and Indiana (Callaway, 1990), and high concentrations of A. triloba occur at the southern borders of these states along the Ohio River (Jacquart et al., 1992; Journal of Heredity, 1917; Zimmerman, 1941). It is therefore noteworthy that alluvial soils of the Ohio River floodplain and its associated terraces bordering the states of Ohio and Indiana are dominated by limestone parent material, underlain by limestone, or underlain by calcareous gravel (e.g., the Huntington–Wheeling and Genesee–Huntsville–Wabash stream bottomland soil associations described by Aandahl et al., 1960). Asimina parviflora (Michx.) Dunal, a related species, is reported to inhabit "lime sinks" within much of the southern U.S. indigenous range of A. triloba (Callaway and Callaway, 1992). Still later observation has revealed the presence of A. triloba as a dominant understory species on a calcareous upland site in central Illinois (Ebinger et al., 1997), and that Ca may be a limiting factor for A. triloba seedling growth in a root-confined (containerized) greenhouse environment (Pomer et al., 2002b). Taken together, these findings support the possibility that improved establishment of A. triloba in the gypsum-amended conditions was, at least in part, related to Ca function in root growth and development.

For trees on gypsum-amended soil, total dry matter accumulation and the net uptake of N, P, and K per tree were at least twice the amount of their nontreated counterparts at the same level of total soil N fertility (expressed as TKN concentration). The relatively poor edaphic establishment environment created by the low-Ca, high-Na conditions may increase residual NO₃ concentration in the soil, which in turn would be subject to various loss mechanisms, particularly leaching. Greater macroelement uptake per tree (Ca-amended conditions) is of practical interest for fertilizer nutrient recovery in orchards (Weinbaum et al., 1992), and for maximizing the environmental benefits of Na-affected agroforestry uses of A. triloba (Robles-Díaz-de-León and Navas-Tudela, 1998). In summary, gypsum supplementation may represent a useful addition to A. triloba orchard management under relatively low saline but high-Na conditions, particularly for increasing growth of trunk and roots during establishment, and for increasing fertilizer nutrient recovery. Intraspecific variation in plant response to Ca has been documented (Snydor, 1962; Snydor and Bradshaw, 1961). Ecotypic variation in genetic composition among marginal populations of A. triloba (e.g., those on the geographic extremities of the species distribution) has also been reported (Huang et al., 1998). Further research is needed to determine whether edaphic differentiation exists among A. triloba populations with respect to their responses to Ca. In addition, more detailed study at the soil : root level could elucidate whether the beneficial effects of gypsum on A. triloba in the Na-affected conditions resulted from improvement of soil physical properties, directly from Ca as nutritional factor, or a combination of both. Such study, for example, could determine the relative importance of Ca fertilization for low-Na A. triloba field plantings.

### Table 3. Macroelement concentrations in roots and combined trunks plus lateral branches, expressed as percentage of dry weight, of excavated A. triloba trees (17 Oct. 1996).³

<table>
<thead>
<tr>
<th>Gypsum rate (t·ha⁻¹)</th>
<th>N</th>
<th>P</th>
<th>K</th>
<th>Ca</th>
<th>Mg</th>
<th>S</th>
<th>N</th>
<th>P</th>
<th>K</th>
<th>Ca</th>
<th>Mg</th>
<th>S</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.0</td>
<td>1.17</td>
<td>0.18</td>
<td>0.75</td>
<td>0.36</td>
<td>0.13</td>
<td>0.19</td>
<td>1.09</td>
<td>0.10</td>
<td>0.50</td>
<td>0.42</td>
<td>0.11</td>
<td>0.07</td>
</tr>
<tr>
<td>7.5</td>
<td>1.17</td>
<td>0.19</td>
<td>0.71</td>
<td>0.39</td>
<td>0.13</td>
<td>0.21</td>
<td>1.04</td>
<td>0.10</td>
<td>0.46</td>
<td>0.42</td>
<td>0.10</td>
<td>0.07</td>
</tr>
<tr>
<td>15.0</td>
<td>1.18</td>
<td>0.19</td>
<td>0.72</td>
<td>0.37</td>
<td>0.12</td>
<td>0.20</td>
<td>1.04</td>
<td>0.10</td>
<td>0.46</td>
<td>0.41</td>
<td>0.10</td>
<td>0.07</td>
</tr>
</tbody>
</table>

³Each value is the mean of six replications.

---

### Literature Cited


U.S. Salinity Laboratory Staff. 1954. Diagnosis and improvement of saline and alkali soils. U.S. Dept. Agr., Wash., D.C.


