



Temporal and Spatial Glucose and Starch Partitioning in Live Oak

Tomas Martinez-Trinidad, W. Todd Watson, Michael A. Arnold,
and Leonardo Lombardini

Abstract. Carbohydrate translocation, which follows anatomical and developmental patterns, is ruled by source-sink relations where energy-containing compounds are moved from sources of production to sinks of utilization. Seasonal carbohydrate concentrations in various tree parts were measured and compared in 10 cm (4 in) trunk diameter live oaks (*Quercus virginiana* P. Miller). Tissue samples from roots, trunks, twigs and leaves were collected from three-year-old field-grown trees on four dates throughout the 2005–2006 seasons. Laboratory analyses of glucose and starch were performed, and values were compared and contrasted according to sample location and time of year. Glucose levels were significantly higher in leaves during the winter ($P \leq 0.001$), while starch concentrations were significantly higher in root and trunk tissues during the spring and winter assessments ($P \leq 0.001$). Carbohydrate concentrations varied among tissues sampled within the tree. This study provides valuable information on the spatial and temporal partitioning of energy reserves, glucose and starch, in live oak so that arborists will have a better understanding of tree vitality, and the effects and environmental impacts of arboricultural treatments.

Key Words. Carbohydrates; *Quercus virginiana*; Source-Sink Relations; Sugar.

Carbohydrates are the principal products of photosynthetic activity and serve as the main energy reserve of trees (Tromp 1983). These products are used by organs where they are produced (e.g., leaves) or are translocated to other organs, a phenomenon controlled by sink-source relations (Allen et al. 2005). Therefore, carbohydrates can be translocated from organs that produce photosynthates (sources) to organs that produce little or no carbohydrates (sinks) where they can be used or stored (Taiz and Zeiger 2006). These sink-source relationships are influenced by tree vitality, nutritional state, environmental conditions, and the developmental stage of plants or tissues (Tschaplinski and Blake 1994; Grulke et al. 2001; Retzlaff et al. 2001). Understanding carbohydrate activity is critical in stressful environments, such as urban forests, where tree health is negatively impacted and environmental stressors are frequent.

Research has highlighted the important role of carbohydrate reserves on a tree's ability to tolerate stressful conditions (Barbaroux et al. 2003). Nonstructural carbohydrates (i.e., starch and soluble sugars) influence the capacity in trees for supporting growth, metabolism, and ultimately their survival (Kaelke and Dawson 2005). Most of the research information about reserve translocation in trees has been explained for young plants grown in greenhouses or natural environments rather than for older trees where carbohydrate production, translocation, utilization, and storage may differ greatly (Tschaplinski and Blake 1994; Gansert and Sprick 1998; Tognetti and Johnson 1999; Domisch et al. 2002). Additionally, much of the recent research concerning carbohydrate partitioning has focused on a single season or short period of time (DeLucia et al. 1998; Retzlaff et al. 2001; Barbaroux et al. 2003).

Trees vary in the allocation or use of carbohydrates stored in tissues. In deciduous trees, roots and trunk serve as the main storage organs during the dormant season, and their reserves are typi-

cally depleted shortly before leaves begin to emerge; conversely, evergreen trees store considerable amounts of starch in leaves and branches (Larcher 1980; Grulke et al. 2001; Retzlaff et al. 2001; Newell et al. 2002). Deciduous trees require extensive carbohydrate storage to maintain the living biomass and cope with stress-inducing factors (Abod and Webster 1991; Gansert and Sprick 1998). For example, white oak (*Quercus alba* L.) rapidly mobilizes and replaces starch reserves during the critical period of canopy generation in the spring (McLaughlin et al. 1980). Conifers accumulate carbohydrates in needles and twigs prior to bud-break and translocate them during the beginning of shoot growth (Ludovici et al. 2002). Some conifers such as Scots pine (*Pinus sylvestris* L.) allocate high percentages of sugar in needles as a response to low soil temperatures (Domisch et al. 2002).

Trees in urban environments are subjected to numerous environmental stressors throughout the year that negatively impact carbohydrate production, utilization, and storage. Live oak (*Quercus virginiana*) is a common species found in urban environments in the southern United States due to the species' elegant canopy shape, adaptability to poor sites, low maintenance requirements, disease resistance, and long life span, which make this species suitable as an ornamental for urban environments (Little 1979; Gilman and Watson 1994). Because a live oak's leaves continue to function throughout the winter until the trees defoliate during budbreak, the species is considered as semi-evergreen. Therefore, research on carbohydrate allocation will help to understand whether the species follows a deciduous or evergreen pattern. Even when under urban conditions, carbohydrate allocation can also be affected when tree organs are modified by human activities such as root pruning, canopy pruning or trunk damage (Harris et al. 2004).

Carbohydrate reserves help to offset low carbohydrate production due to stressful conditions or high demand. Differences in

the allocation of carbohydrates to storage tissues could arise from varied requirements of different organs, different needs during growth, or maintenance respiration required among different species (Dean 2001; Barbaroux et al. 2003). Variation in starch concentrations could indicate different rates of production, demand, or shifts in allocation (Ludovici et al. 2002). In trees, the continuous pathways of transport and storage of previous-year assimilates are essential for subsequent growth processes, so the coordination and interrelations of morphogenic and photosynthetic processes are very important (Kaipiainen and Sofronova 2003). Carbohydrates can also be used under stress conditions as a precursor for secondary compounds used to resist biotic or abiotic stress (Webb 1981; Renaud and Mauffette 1991). Understanding carbohydrate partitioning in live oaks can help in understanding the translocation of the main carbohydrate, glucose and starch in semi-evergreen species as well as the relationships between phenological stages and movement and utilization of available energy resources.

The objective of this research was to study the non-structural carbohydrate content, glucose and starch, in tree roots, trunks, twigs, and leaves of large, field-grown live oaks as well as to determine the impacts of seasonal influences on carbohydrate concentrations and utilization.

MATERIAL AND METHODS

Plant Material

Five, field-grown live oaks, about 10 cm (4 in) trunk diameter measured 30 cm (12 in) above ground, were randomly selected from within a nursery at Monaville, TX (29°57'1.59"N, 96°3'28.73"W). Trees selected in the 2005-year were planted in 1999 at 5 m (16.4 ft) spacing, and grown under similar conditions. The soil was a deep, moderately well drained, slowly permeable Lake Charles clay, and trees were annually fertilized (20-20-20, water soluble tree fertilizer).

Tissue samples from roots, trunk, and canopy were collected from opposing sides of the tree (across and between rows of trees) corresponding to the four cardinal points. Root samples consisted of 4 mm (0.16 in) diameter increment cores from the buttress (woody) roots [2 cm (0.79 in) from the base of the trunk]. Tissues from the trunk, cores 4 mm diameter, were collected using an increment hammer (Haglof®; Langsele, Sweden) at 1.3 m (4.3 ft) height from the ground. Increment cores were approximately 100 mm (3.94 in) in length. Canopy samples consisted of five 5 cm (2 in) long twigs with leaves, which were randomly collected from the lower two-thirds of the canopy (McLaughlin et al. 1980). Samples from the different parts of the trees were collected within two hours in July 2005, September 2005, January 2006, and March 2006.

Samples were stored on blue ice (Rubbermaid®, Fairlawn, OH) immediately after collection in the field, transported to the lab within two hours, and oven-dried at 80 °C (176 °F) until weights stabilized (approximately 48 hours). After drying, samples were ground and stored in plastic bottles at -20 °C (-4 °F) until the carbohydrate concentrations were analyzed (Kolb and McCormick 1991).

Carbohydrate Analysis

Glucose and starch concentrations were determined for each sample using Sigma® GAGO-20 reagents (Sigma, St. Louis, MO). Glucose was extracted from tissues using methanol:chloroform:water (MCW, 12:5:3, v/v/v) solution, and 0.5 mL (0.02 fl oz) of the su-

pernatant from the extract or glucose standards was mixed with 5 ml (0.17 fl oz) of anthrone reagent (Jaenicke and Thiong'o 1999). Absorbance of samples and standards were read at 625 nm within 30 minutes using a spectrophotometer (Spectronic 20, Baush & Lomb, Rochester, NY). Glucose concentrations were calculated through standard linear regressions and expressed as mg per g of dry weight. Starch content was determined by enzymatic conversion of starch to glucose. In the remaining pellet after glucose extractions, amyloglucosidase, an enzyme responsible for the conversion of starch to glucose, was used. Standards were prepared using potato starch, and the samples were read at 540 nm within 30 minutes. Starch content was calculated through standard linear regressions and expressed as mg of glucose per g of dry weight. (Haissig and Dickson 1979; Renaud and Mauffette 1991).

Data Analysis

An analysis of variance was conducted on data including in the model cardinal orientation, type of tissue, and time of year. Results from samples collected from opposite side of the tree (North-South, and West-East) were pooled in order to compare differences among and between rows in the nursery. The data was analyzed with the procedure General Linear Model (GLM) using the Statistical Package for the Social Sciences (SPSS, v. 13) for Windows (SPSS, Chicago, IL). Comparisons among tissues and within orientations were performed using Fisher's Least Significant Differences (LSD) to determine differences in carbohydrate concentration in trees.

RESULTS AND DISCUSSION

Statistical analysis revealed no main or interactive effect ($P > 0.5$) of cardinal orientation. When data from opposing orientations (N-S and E-W) were pooled, no differences were found (data not shown). These results indicate that future research involving tissue sampling for carbohydrates may not be impacted by cardinal direction when sampling similar tissues at similar heights. However, carbohydrate reserve concentrations varied from base to the top in the trunk of sessile oak (*Quercus petraea* L.) and beech (*Fagus sylvatica* L.) (Babaroux et al. 2003).

Differences in glucose concentrations among tissues varied ($P < 0.001$) across seasons (Figure 1). Glucose content was higher in leaves than in other tissues tested through the year, except in September when there were no significant differences in glucose

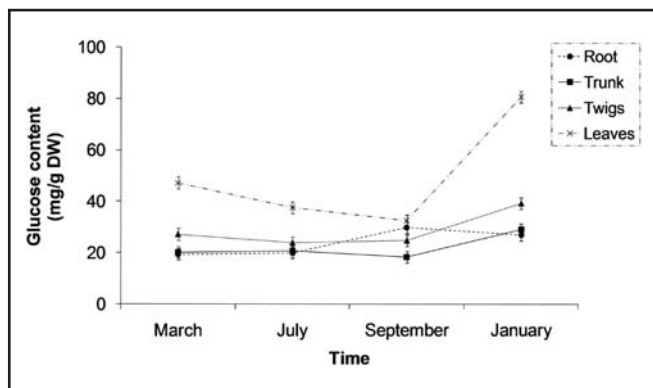


Figure 1. Glucose concentrations (mg/g dry weight) measured in live oak samples collected at four different dates. Bars represent the SE of the mean.

levels in leaves and roots. Although leaf glucose levels decreased in September, they were still higher than glucose levels in trunk and twigs. To the contrary, species such as pecan (*Carya illinoensis* (Wangenh) K. Koch) or ponderosa pine (*Pinus ponderosa* Laws.) showed constant concentration of glucose during the growing season in different tissues of the tree (Gulke et al. 2001; Kim and Wetzstein 2005). Foliar glucose levels decreased in July and September, possibly due to the developmental stage of leaves (Figure 1). The glucose fraction in roots reached the highest concentration in the fall which may likely indicate a high translocation of sugars to roots for storage purposes. Root glucose levels were lower in the March–July period, which corresponds to the time when the roots had the highest levels of starch (Figure 2). The decrease in glucose content in leaves during fall can be a result of the high translocation and replacement of carbohydrates from source to sinks or storing organs (McLaughlin et al. 1980).

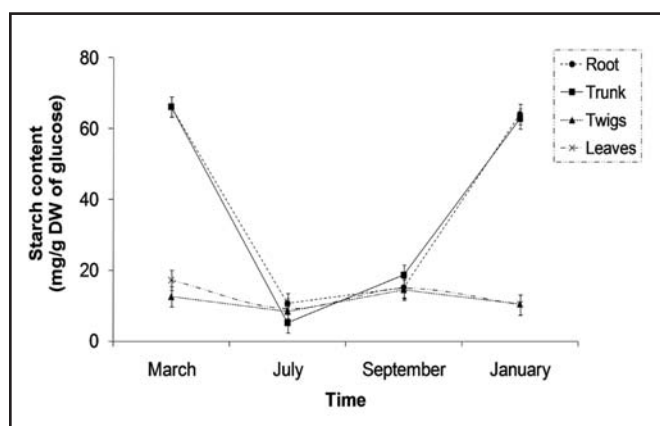


Figure 2. Starch concentrations (mg/g DW of glucose) measured in live oak samples collected at four different dates. Bars represent the SE of the mean.

Glucose concentrations in leaves might increase in winter because live oaks maintain live leaves throughout the winter as a response to environmental factors. Ludovici et al. (2002) also found a two-to-three-fold increase in winter glucose concentrations when compared to summer levels in needles of loblolly pine (*Pinus taeda* L.). The high levels of glucose in leaves during the winter assessment can be caused as a response to the low temperatures [below 0 °C (32 °F)] that occurred before the sample collection. Conversion of starch to sugar is a physiological manifestation of cold hardiness in trees (Levitt 1980; Nguyen et al. 1990).

A significant interaction ($P < 0.001$) was also found for starch among time of year and tissues sampled. The highest starch concentrations were measured in roots and trunks during spring and winter (Figure 2), thus confirming their role in facilitating carbohydrate storage (Allen et al. 2005). Similar results were found for trunks and roots of sessile oak which had higher quantities of carbohydrate reserves in autumn than in late summer (Barbaroux et al. 2003). Trunk tissues showed a depletion of starch from spring to the summer (Figure 2). Gansert and Sprick (1998) identified starch disappearance during the summer in beech. Low concentrations of starch were also found during the spring in Scots pine (Domisch et al. 2002).

Starch concentrations found in roots and trunk (Figure 2) emphasize the importance of these two organs as carbohydrate reservoirs during the dormant season. These storage tissues have an

indispensable role during the initiation of vegetative growth in the spring. The reduction of starch levels observed in the summer is likely a consequence of growth and high night temperatures and is associated with the hydrolyzation of stored carbohydrates for growth and maintenance (Kaipainen and Sofronova 2003; Pallardy 2008).

Even when the interaction of type of tissue and time of year was significant, the annual mean carbohydrate concentrations varied among different tissues in trees (data not shown). Glucose levels in leaves were about double than those in twig, root, and trunk tissues. For starch, an inverse pattern was found where root and trunk tissues had more than three times the starch content than foliar tissues. This overall concentration pattern corroborates source-sink mechanisms for carbohydrate partitioning described in other studies (Kaipainen and Sofronova 2003; Kaelke and Dawson 2005; Taiz and Zeiger 2006). Dean (2001) indicated that root allocation can be affected by stem competition because stems precede roots on the chain of carbohydrate sinks.

For future studies aimed at monitoring carbohydrate levels in trees, additional factors must be considered. For example, twigs exhibited less variation compared with other tissues analyzed and were easy to collect any time of year without causing considerable severe injury to the trees. Leaves are only available during the growing season and it requires being careful to obtain a homogenous sample during the dormant season. In the case of sampling roots and trunks, the tree sustains more damage that may disrupt physiological functions. In addition, fine root sampling can be complicated due to the partial loss of the sample during the washing process (Kaipainen and Sofronova 2003). Changes depending on species have also been reported for carbohydrate partitioning (Newell et al. 2002). Based on the results of this study, trunk, root, twig, or leaf tissue, and can be used to monitor glucose and starch levels throughout the year; however, the seasonal variation in different parts of the tree and the different species should be taken into consideration.

LITERATURE CITED

- Abod, S.A. and A.D. Webster. 1991. Carbohydrates and their effects on growth and establishment of *Tilia* and *Betula*: I. Seasonal changes in soluble and insoluble carbohydrates. *Journal of Horticulture Science* 66:235–246.
- Allen, M.T., P. Prusinkiewicz, and T.M. Dejong. 2005. Using L-systems for modeling source-sink interactions, architecture and physiology of growing trees: the L-PEACH model. *New Phytologist* 166:869–880.
- Barbaroux C., N. Breda, and E. Dufrene. 2003. Distribution of above-ground and below-ground carbohydrate reserves in adult trees of two contrasting broad-leaved species (*Quercus petraea* and *Fagus sylvatica*). *New Phytologist* 157:605–615.
- Dean, T.J. 2001. Potential effect of stand structure on belowground allocation. *Forestry Science*. 47:69–76.
- DeLucia, E.H., T.W. Sipe, J. Herrick, and H. Maherali. 1998. Sapling biomass allocation and growth in the understory of a deciduous hardwood forest. *American Journal of Botany* 85:955–963.
- Domisch, T., L. Finer, and T. Lehto. 2002. Growth, carbohydrate and nutrient allocation of Scots pine seedlings after exposure to simulated low soil temperature in spring. *Plant Soil* 246:75–86.
- Gansert, D. and W. Sprick. 1998. Storage and mobilization of nonstructural carbohydrates and biomass development of beech seedlings (*Fagus sylvatica* L.) under different light regimes. *Trees* 12:247–257.
- Gilman, E.F. and D.G. Watson. 1994. *Quercus virginiana*: southern live oak. U.S. Dept. Agriculture Forest Service Fact Sheet. ST-564.

- Grulke, N.E., C.P. Andersen, and W.E. Hogsett. 2001. Seasonal changes in above-and belowground carbohydrate concentrations of ponderosa pine along a pollution gradient. *Tree Physiology* 21:173–181.
- Haissig, B.E. and R.E. Dickson. 1979. Starch measurement in plant tissue using enzymatic hydrolysis. *Physiologia Plantarum*. 47:151–157.
- Harris, R.W., J.R. Clark, and N.P. Matheny. 2004. *Arboriculture: integrated management of landscape trees, shrubs, and vines*. Prentice Hall, Upper Saddle River, NJ.
- Jaenicke, H. and M. Thiong'o. 1999. Comparison of two methods for the routine extraction and quantification of non-structural carbohydrates in tree tissues. *Journal Agriculture Science and Technology* 2:46–55.
- Kaelke, C.M. and J.O. Dawson. 2005. The accretion of nonstructural carbohydrates changes seasonally in *Alnus incana* ssp. *rugosa* in accord with tissue type, growth, N allocation, and root hypoxia. *Symbiosis* 39:61–66.
- Kaipainen, L.K. and G.I. Sofronova. 2003. The role of the transport system in the control of the source-sink relations in *Pinus sylvestris*. *Russian Journal of Plant Physiology* 50:125–132.
- Kim, T. and H.Y. Wetzstein. 2005. Seasonal fluctuations in nutrients and carbohydrates in pecan leaves and stems. *Journal of Horticulture Science. Biotech.* 80:681–688.
- Kolb, T.E. and L.H. McCormick. 1991. Relationship between total non-structural carbohydrate concentration and root diameter in sugar maple. *Forestry Science* 37:343–346.
- Larcher, W. 1980. *Physiological plant ecology*. 2nd Ed. Springer-Verlag, New York.
- Levitt, J. 1980. *Responses of plants to environmental stress*, Volume 1. Academic Press, New York.
- Little, E.L. 1979. Checklist of United States trees (native and naturalized). U.S. Dept. Agriculture Forest Service Handbook 541 Washington, D.C.
- Ludovici, K.H., H.L. Allen, T.J. Albaugh, and P.M. Dougherty. 2002. The influence of nutrient and water availability on carbohydrate storage in loblolly pine. *Forest Ecology Management* 159:261–270.
- McLaughlin, S.B., R.K. McConathy, R. L. Barnes, and W.T. Edwards. 1980. Seasonal changes in energy allocation by white oak (*Quercus alba*). *Canadian Journal Forestry Research* 10:379–388.
- Newell, E.A., S.S. Mulkey, S.J. Wright. 2002. Seasonal patterns of carbohydrates storage in four tropical tree species. *Oecologia* 131:333–342.
- Nguyen, P.V., D.I. Dickmann, K.S. Pregitzer, and R. Hendrick. 1990. Late-season changes in allocation of starch and sugar to shoots, coarse roots, and fine roots in two hybrid poplar clones. *Tree Physiology* 7:95–105.
- Pallardy, S.G. 2008. *Physiology of woody plants*. 3rd. Ed. Academic Press, New York.
- Renaud, J.P. and Y. Mauffette. 1991. The relationship of crown dieback with carbohydrate content and growth of sugar maple (*Acer saccharum*). *Canadian Journal Forestry Research* 21:1111–1118.
- Retzlaff, W.A., J.A. Handest, D.M. O'Malley, S.E. McKeand, and M.A. Topa. 2001. Whole-tree biomass and carbon allocation of juvenile trees of loblolly pine (*Pinus taeda*): influence of genetics and fertilization. *Canadian Journal Forestry Research* 31:960–970.
- Taiz, L. and E. Zeigler. 2006. *Plant physiology*. 4th Ed. Sinauer Associates Inc., Sunderland, MA.
- Tognetti, R. and J.D. Johnson. 1999. The effect of elevated atmospheric CO₂ concentration and nutrient supply on gas exchange, carbohydrates and foliar phenolic concentration in live oak (*Quercus virginiana* Mill) seedlings. *Ann. Forestry Science* 56:379–389.
- Tromp, J. 1983. Nutrient reserves in roots of fruit trees in particular carbohydrates and nitrogen. *Plant Soil* 71:401–413.
- Tschaplinski, T.J. and T.J. Blake. 1994. Carbohydrate mobilization following shoot defoliation and decapitation in hybrid poplar. *Tree Physiology* 14:141–151.
- Webb, W.L. 1981. Relation of starch content to conifer mortality and growth loss after defoliation by the Douglas-fir tussock moth. *Forestry Science* 27:224–232.

Acknowledgments. We thank Environmental Design Inc. for supporting this research.

Tomas Martinez-Trinidad (corresponding author)
Colegio de Postgraduados – Programa Forestal
Km. 36.5 Carr. Mex-Tex. Montecillo
Texcoco, Edo. de Mexico. 56230. Mexico
tomtz@colpos.mx

W. Todd Watson
Department of Ecosystem Science & Management
Texas A&M University
College Station, TX, U.S.
t-watson@tamu.edu

Michael A. Arnold
Department of Horticultural Sciences
Texas A&M University
College Station, TX 77843, U.S.
ma-arnold@tamu.edu

Leonardo Lombardini
Department of Horticultural Sciences
Texas A&M University
College Station, TX 77843, U.S.
l-lombardini@tamu.edu

Résumé. La translocation des hydrates de carbone qui suit des patrons de développement et d'anatomie spécifiques est régie par des relations de source-vide où les molécules porteuses d'énergie sont transportés des zones source de production vers les zones d'utilisation qui sont en besoin. Les concentrations saisonnières en hydrates de carbone de diverses portions de l'arbre ont été mesurées et comparées parmi des chênes verts (*Quercus virginiana* P. Miller) de 10 cm de diamètre de tronc. Des échantillons de tissus provenant de racines, de troncs, de tiges et de feuilles ont été recueillis auprès d'arbres en champ de trois ans à quatre moments différents en 2005 et 2006. Des analyses en laboratoire de glucose et d'amidon ont été menées et les valeurs comparées entre elles en fonction de la localisation et de la période de l'année. Les niveaux de glucose étaient significativement plus élevés dans les feuilles durant l'hiver ($P \leq 0,001$), alors que les concentrations en amidon étaient significativement plus élevées dans les tissus de racines et de troncs au printemps et en hiver ($P \leq 0,001$). Les concentrations en hydrates de carbone variaient parmi les tissus au sein d'un même arbre. Cette étude fournit de l'information précieuse sur la répartition spatiale temporelle des réserves d'énergie, glucose et amidon, chez le chêne vert ce qui permet aux arboriculteurs d'avoir une meilleure compréhension de la vitalité de l'arbre ainsi que des effets et des impacts environnementaux des traitements arboricoles.

Zusammenfassung. Die Kohlenhydrat-Translokation, welche nach anatomischen und entwicklungsentsprechenden Strukturen abläuft, ist beherrscht durch Source-sink-Beziehungen, wo die energiehaltigen Bestandteile von der Quelle der Produktion zur Stelle des Einsatzes transportiert werden. Bei jungen Lebenseichen mit 10 cm Stammdurchm. wurden saisonale Kohlenhydratkonzentrationen in verschiedenen Baumteilen gemessen und verglichen. In der Saison 2005 bis 2006 wurden viermal an dreijährigen Bäumen Gewebeproben aus Wurzeln, Stamm, Zweigen und Blättern entnommen. Im Labor wurde der Glukose- und Stärkegehalt bestimmt, die Werte verglichen und entsprechend ihrer Entnahmestelle

und Jahreszeit in Vergleich gesetzt. Die Glukosegehalte waren in den Blättern im Winter deutlich höher, während die Stärkekonzentrationen im Winter und Frühjahr in den Wurzeln und Stammgewebe deutlich höher waren. Die Kohlenhydratkonzentrationen variierten innerhalb des Baumes. Diese Studie liefert wertvolle Informationen über die räumliche und temporäre Partitionierung von Energiereserven, Glukose und Stärke, in Lebenseichen, so dass Arboristen nun ein besseres Verständnis für die Vitalität und die Auswirkungen von Pflegemaßnahmen auf die Bäume bekommen.

Resumen. La translocación de carbohidratos, la cual sigue patrones de desarrollo anatómicos, está regida por las relaciones fuente-recipiente donde los compuestos energéticos son movidos de fuentes de producción a recipientes de utilización. Las concentraciones estacionales de carbohidratos en varias partes del árbol fueron medidas y comparadas en encinos siempreverdes (*Quercus virginiana* P. Miller) con diámetros del tronco de 10 cm (4 pulg). Se colectaron muestras de tejido de raíces, troncos, brotes y hojas de árboles de campo de tres años en cuatro fechas, a través de las estaciones, en 2005–2006. Se realizaron análisis de laboratorio de glucosa y almidón y los valores fueron comparados y contrastados de acuerdo a la localización de la muestra y época del año. Los niveles de glucosa fueron significativamente más altos en las hojas durante el invierno ($P \leq 0.001$), mientras que las concentraciones de almidón fueron significativamente más altas en los tejidos de las raíces y el tronco durante las mediciones de primavera e invierno ($P \leq 0.001$). Las concentraciones de carbohidratos variaron entre los tejidos muestreados dentro del árbol. Este estudio proporciona información valiosa sobre la distribución espacial y temporal de las reservas de energía, glucosa y almidón en encinos, por lo que los arboristas tendrán un mejor entendimiento de la vitalidad del árbol y los efectos e impactos ambientales de los tratamientos arboriculturales.