

# Modeling Actual Evapotranspiration of *Viburnum odoratissimum* during Production from Rooted Cuttings to Market Size Plants in 11.4-L Containers

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**Abstract.** Rooted cuttings of *Viburnum odoratissimum* were grown outdoors to market size in 11.4-L containers. Actual evapotranspiration ( $ET_A$ ) of nine plants was determined daily as was evaporation from three control containers shaded with plastic foliage to mimic plant growth. The first 60 d after transplanting, substrate evaporation accounted for most of  $ET_A$ . Substrate evaporation was generally constant the first 160 days before declining, but still remained  $\approx 160$  mL/day through harvest at market size.  $ET_A$  increased with growth and generally followed variations in reference evapotranspiration ( $ET_o$ ). Mean  $ET_A$  during most of the production cycle was less than 600 mL/day (11.8 mm based on upper container surface area). With the spring growth flush, mean  $ET_A$  reached 1.3 L/day as plants achieved market size. Mean cumulative  $ET_A$  to produce 90% of measured plants to market size was 155 L or 3.1-m depth per plant based on container surface area. Water need indices, similar to crop coefficients, were highly correlated with percent canopy closure (%Closure) using an exponential decay equation. When overlain with previous similar data for *Ligustrum japonicum*, the correlation for the combined data set had an  $r^2 = 0.843$ . This suggests that the %Closure model may provide a method for  $ET_o$ -based irrigation of woody shrub species based on canopy size and spacing.

Container production of woody ornamentals requires frequent and abundant irrigation. In Florida, irrigation up to 2.9 m/year was common in the mid-1970s in addition to an annual mean rainfall of 1.1 m (Harrison, 1976). In 1992, maximum allowed rates of 2.29 m of supplemental irrigation were imposed by Florida's Water Management Districts. In 2003, this maximum was reduced 22% for nurseries where competition for potable groundwater with expanding urban centers occurred. With further restrictions probable, it becomes imperative to know how much irrigation is required to produce container plants and to develop ways nurseries can become more precise in their irrigation applications.

In Florida, overhead irrigation of nurseries has been restricted during most of the daylight hours since the mid-1990s (Southwest Florida Water Management District, 2009). To cope with this imposition, along with decreased labor pools, irrigation at most nurseries queried occurs at night and is controlled by time clock systems (Beeson, personal communication). Most irrigation

managers are unaware of day-to-day changes in reference evapotranspiration ( $ET_o$ ), a value that quantifies evaporation potential of the immediate microclimate, and corresponding changes in actual evapotranspiration ( $ET_A$ ) of their plants. During the summer, thunderstorm rainfall is common after employees depart for the day, often with time clocks preset. Rain shutoff devices are required, but they do not function accurately or consistently. Because irrigation in many nurseries is already centrally controlled and using electric valves, implementation of a simple and accurate algorithm relating daily environmental conditions to container irrigation needs could improve plant growth and conserve water.

Attempts to develop such an irrigation model for container-grown woody shrubs have been limited. Knox (1989) reported relatively high correlations between pan evaporation, a growth index, and  $ET_A$  for five woody ornamentals during a year's production cycle. However, a working model was not reported. Fitzpatrick (1983a, 1983b) reported water consumption for tropical ornamentals along with monthly  $ET_o$  derived from the older Thornthwaite (Thornthwaite, 1948) equation. Irrigation was based on a previously developed model (Fitzpatrick, 1980) but was empirically dependent on a specific growth rate and monthly estimates of  $ET_o$

based on historical data and thus had limited usefulness.

In more recent years, modifications of the Penman-Monteith equation (Allen et al., 1998; Walter et al., 2002) have served as the basis to make daily estimates of  $ET_o$  and calculations of corresponding crop coefficients (Kc). Crop coefficients are unitless values that relate  $ET_A$  to  $ET_o$  and are determined empirically yet robust in application. For container-grown plants,  $ET_A$  is most commonly calculated as changes in mass of the container-plant system determined by weighing. However  $ET_o$  is calculated as a depth. To calculate a Kc,  $ET_A$  first must be converted to a depth of water by dividing by an area. Burger et al. (1987) used the container top diameter for calculating the surface area to convert  $ET_A$  to a depth for 22 woody ornamental species growing in 3.8-L containers. At market size, Kc ranged from 1.1 to 5.1, depending on species and container spacing. High Kc values, relative to agronomic crops that rarely exceed 1.3 (Doorenbos and Pruitt, 1977), were justified by large canopy to container surface area ratios (Burger et al., 1987). Differences in Kc values between geographically distant locations were minimum. Niu et al. (2006) used the same methodology to calculate Kc for five woody species in western Texas in two container sizes with different volumes but identical top diameters. Plants and containers in this study were larger than those of Burger et al. (1987). Kc values based on container top diameter ranged from 0.87 to 4.37, similar to those reported previously for smaller plants (Burger et al., 1987; Regan, 1997). Whereas water use per unit leaf area was similar between container sizes within species, Kc values of different container sizes within species varied significantly as a result of differences in plant size. This underscores a problem with calculating Kc based on container top diameter; Kc is dependent on both container and plant size and varies with time (Beeson, 1996; Niu et al., 2006). Additionally, container top diameter varies depending on manufacturer and model. Recently Bacci et al. (2008) reported on a method to adjust Kc for plant growth by combining Kc as derived by Burger et al. (1987) with measurements of substrate moisture content derived from tensiometers. Kc is adjusted based on a sliding 7-d average of substrate moisture content. The system offers two concurrent, mostly independent methods of estimating daily irrigation requirements. This system relies on accurate measurements of soil matric potential, a parameter subject to measurement errors resulting from sensor placement, root growth, and especially substrate composition and time.

In 1996 Beeson (1996) reported linear relationships between  $ET_o$  and  $ET_A$  based on projected canopy area (PCA) and container substrate surface area. PCA was calculated by multiplying the widest canopy width by the width perpendicular to the widest width. Correlations ranged from 0.533 to 0.695 for well-irrigated treatments when  $ET_A$  depth

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was based on PCA. Correlations were also significant but lower when  $ET_A$  depth was based on container surface area ( $r^2 < 0.30$ ). The Kc's for these *Ligustrum japonicum* Thunb. ranged from 1.28 to 6.12 when based on container surface area, similar to those reported by others (Burger et al., 1987; Niu et al., 2006; Regan, 1997). However Kc ranged from 0.50 to 0.63 when based on PCA, which is within the range of Kc reported for agronomic and tree crops. For all of these studies, data collection occurred over a single growing season.

Schuch and Burger (1997) were the first to attempt modeling Kc for multiyear woody ornamental crops using ETo. Over a 20-month production cycle, Kc ranged from 1.0 to 2.5 for woody ornamentals grown in 15.6-L containers based on container top diameter. To account for quiescent periods during winter, a Fourier curve transformation was used. In 2004, a model was presented for *L. japonicum* relating  $ET_A$  to ETo, plant canopy size, and spacing between containers (Beeson, 2004). The model, called the Percent Canopy Closure model, was based on daily ETo and  $ET_A$  determined by weighing lysimetry for *L. japonicum* during a year of production from rooted cuttings to market size plants. In 2005, an algorithm derived from the model was used to control irrigation of *L. japonicum* from rooted cuttings to market size plants (Beeson and Brooks, 2008). The algorithm was successful in producing a crop 3 weeks earlier and with less irrigation than the manually adjusted control that was targeted for 1.8 m of irrigation annually.

The primary objective of the research presented here was to quantify cumulative  $ET_A$  of *Viburnum odoratissimum* Ker Gawl (viburnum) to expand the database of woody shrub water requirements during production. The second objective was to analyze the data using the Percent Canopy Closure model and compare results with those obtained earlier with *L. japonicum*.

## Materials and Methods

On 3 Mar. 2005, rooted cuttings of *Viburnum odoratissimum* produced in 5.7-cm peat pots (70000041; Jiffy Products of American, Lorain, OH) were transplanted into 11.4-L black polyethylene containers (C1200; Nursery Supply, Chambersburg, PA) at the Mid-Florida Research and Education Center in Apopka, FL. Substrate was commercially prepared (Florida Potting Soil, Inc., Apopka, FL) and consisted of 55% pine bark fines:36% Nupeat:9% sand amended with 2.9 kg·m<sup>-3</sup> dolomite limestone and 0.86 kg·m<sup>-3</sup> micro-nutrients (Micro-Max; Scotts Company Inc., Marysville, OH). NuPeat is a mixture of equal parts of composted hardwood bark, composted yard waste, and Florida sedge peat that passes through a 12.5-mm screen. Containers were placed on black polyethylene ground cloth in a square arrangement within a production area measuring 9.2 m × 9.8 m and irrigated at 1200 HR as needed with overhead

impact sprinklers situated at each corner. Before container placement, sprinkler heads were adjusted to achieve a Christiansen Uniformity Coefficient of 0.85 or better (Haman et al., 1977). Application rate was also calculated at that time.

Nine plants were selected and placed in suspension lysimeters for 30-min recording of individual container-plant mass. Lysimeters consisted of 22.7-kg load cells (SSM-AJ-50; Interface Inc., Scottsdale, AZ) suspended from 2.2-m tripods from which a basket was hung that suspended a container ≈25 mm above the ground surface. Tops of lysimeter containers were less than 4 cm above surrounding containers. Load cells were read using a CR10X data logger and AM416 and AM-32 multiplexers (Campbell Scientific, Inc., Logan, UT). Lysimeters were placed in the middle of three banks of plants to prevent edge effects in a completely randomized design. Each bank was five rows wide and 19 plants long. Three additional lysimeters suspending substrate-filled containers without plants were included within the production area to measure evaporation. Plastic foliage was added to these as needed to mimic canopy shading of substrate surfaces that occurred in containers of measured plants and did not interfere with substrate rehydration. Containers were spaced on 0.44-m centers at the inception of the study and not re-spaced. Daily  $ET_A$  for live plants and evaporation for controls were calculated by the data logger based on changes in container-plant mass between 0600 HR and 2200 HR. For days with rain, calculations involved manual separation of  $ET_A$  from rain-induced changes in mass for each lysimeter as detailed by Beeson (2006).

Daily reference evapotranspiration (ETo) was calculated using Campbell Scientific's Application Note 4. Temperature and relative humidity were recorded using a CS-215 sensor (Campbell Scientific). Wind was measured with an anemometer (014A; Met One, Grants Pass, OR) at 2 m above ground level. Solar radiation was measured with a pyranometer (LI-200X; LI-COR Inc, Lincoln, NE). ETo was calculated from data collected within 50 m of the experimental site.

For the first 3 weeks after transplanting, all plants were irrigated with 6 mm of overhead irrigation daily to establish the plants. Thereafter, irrigation treatments were controlled by the lysimeter system. Irrigation of the production area was based on maximum  $ET_A$  of lysimeter plants multiplied by a canopy shedding/irrigation uniformity factor set at 1.0 initially. This was increased in late summer to 1.2 to account for canopy shedding of overhead irrigation once canopies extended beyond container diameter (Beeson and Yeager, 2003). After the first 3 weeks, irrigation depths less than 6.3 mm were deferred to the next day and added to that day's irrigation application.

Each container was top-dressed with 55 g of a controlled-release fertilizer (18N–2.6P–9.9K; 18-6-12 Polyon, 8 to 9 months; Harold's Fertilizer Inc, Clermont, FL) on 8 Mar.

2005 and was treated with pre-emergence herbicide (Ornamental Herbicide II; Scotts Co.). On 4 Apr. 2005, each container was given ≈150 mL of a 300 ppm nitrogen liquid fertilizer (Peter's 20-20-20; Scotts Co.) solution by hand. On 23 Feb. 2006, each container received 34 g of 14N–6P–11.6K Osmocote (14-14-14; Scotts Co.).

Canopy measurements of widest width, width perpendicular to widest width, and average height were recorded on each lysimeter plant and four adjacent border plants generally every 3 weeks beginning shortly after transplanting. Canopy measurements were also recorded for border plants of the three control containers. The two widths were multiplied to calculate PCA for each plant. On 25 Mar. 2006, greater than 90% of measured plants had achieved market size based on Florida Grades and Standards (DACS, 1994). For this species and container size, minimum market size was an average height of 0.60 m and average width of 0.45 m.

Daily  $ET_A$  and evaporation were analyzed as repeated measures using GLM in SAS (Version 9.2; SAS Institute Inc., Cary, NC) with container water loss ( $ET_A$  and evaporation) as the main plot and time as the subplot (Snedecor and Cochran, 1980). *t* tests ( $P < 0.05$ ) were used to compare between variables for each day.

The %Closure was calculated for each lysimeter plant for each measurement date by adding the mean of half the PCA of each border plant to half the PCA of the respective lysimeter plant and then dividing by allocated bed area per plant (1972 cm<sup>2</sup>). Allocated bed area was essentially the square of the distance on-center between plants. With growth, canopies filled the space between containers. Because containers were not re-spaced, canopies overlapped each other as individual shoots of each plant expanded outward and up. Because %Closure was calculated on a fixed allocated bed area, this overlap resulted in calculation of %Closure greater than 100%.

For modeling, only data for a 7-d period centered on each growth measurement date was used. Daily  $ET_A$  (cm<sup>3</sup>) of each lysimeter plant was converted to a depth by dividing by its PCA (cm<sup>2</sup>).  $ET_A$  depth (cm) was then divided by the corresponding ETo (cm) and averaged over the 7 d to estimate a water needs index (WNI; Beeson and Brooks, 2008) for each lysimeter plant at each measurement date. WNI is similar to a Kc but does not meet the constraints of a large uniform fetch as defined for Kc (Doorenbos and Pruitt, 1977). The likelihood of a uniform fetch, an expansive extent of uniform vegetation of the same species, in production nurseries is rare. Hence, coefficients derived for ornamental nursery plants, by definition, cannot be called crop coefficients. WNI was coined to provide a name for coefficients derived from ornamental plant research. WNI values of the nine lysimeter replicates for each date were plotted against their respective %Closure. The plot was fitted to a three-parameter exponential decay curve as

reported by Beeson (2004) using SigmaPlot (Version 10; SPSS Science, Chicago, IL). Data from the viburnum were combined with similar previous data from *Ligustrum japonicum* and the plot fitted to the same exponential decay curve using SigmaPlot.

## Results and Discussion

As was found with *L. japonicum* in 2001 (Beeson, 2004), for the first 56 d after transplanting, evaporation from substrate surfaces accounted for most of the daily water loss (Fig. 1). During this period, there were no differences between  $ET_A$  and evaporation. Between then and the next 34 d,  $ET_A$  exceeded evaporation only 9 d. After the first 90 d,  $ET_A$  exceeded evaporation of artificially shaded containers each day except for 10 d. Eight of these were associated with rain amounts greater than 8.8 mm. The other 2 d were in mid-December when  $ETo$  was less than 0.44 mm and rain occurred. Daily water loss ranged from 50 to 300 mL (1.0 to 6.3 mm based on substrate surface area) from transplanting until approximately Day 160 (90 d after transplanting). Low water losses from Day 98 to 108 were the result of malfunctioning irrigation solenoids. Irrigation was not supplied; thus, evaporation and transpiration were diminished these days. Viburnum transplanted in early spring would require irrigations of 6.3 mm or less daily for the first 3 months given similar  $ETo$  conditions (Fig. 2). This was similar to that reported for *L. japonicum* in 2001 (Beeson, 2004).

Increases in  $ET_A$  began in late June (approximately Day 175) with  $ET_A$  being generally 50% more than evaporation as a result of increases in shoot size represented by mean canopy width (Fig. 3). Daily  $ETo$  was generally less than 1 month previous (Fig. 2). In Florida, daily  $ETo$  peaks in May to early June at the end of the dry season. Peak  $ETo$  occurs as a result of low humidity and little cloud cover combined with solar radiation nearing its peak. Generally the wet season begins the second week of June and continues until early October. The wet season is associated with midday to afternoon cloud cover and high relative humidity that generally limits daily  $ETo$  to 5 mm or less.

Evaporation from artificially shaded containers was relatively constant at 200 mL (4 mm) until early October (Day 275). At this point, mean canopy width of lysimeter plants was 33.5 cm with a %Closure of 51% (Fig. 3). Thereafter, evaporation declined slowly to a baseline of 160 mL (3.1 mm) by late November (Day 325) and remained relatively constant until plants obtained market size. During these last 5 months of near constant substrate evaporation,  $ETo$  varied between 1.2 to 4.6 mm/day (Fig. 2). This suggests that during this period, substrate evaporation was not coupled to the environment (Noble, 1999). In previous research with viburnum,  $ET_A$  declined 40% when %Closure increased from 67% to 100% attributed principally to reductions in whole canopy transpiration caused by decoupling of lower leaves (Beeson, 2010). By

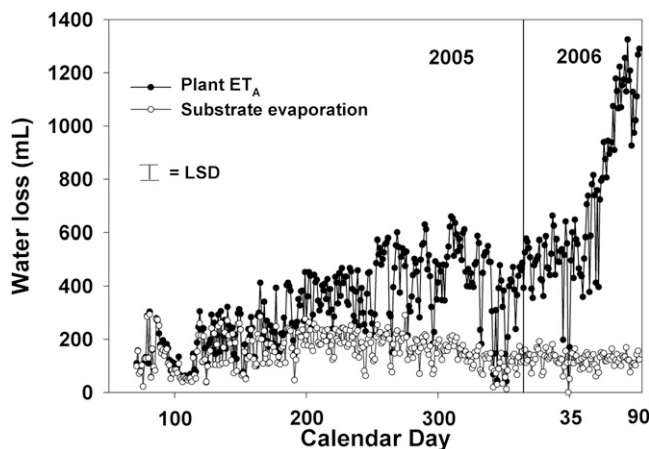


Fig. 1. Mean daily water loss through actual evapotranspiration ( $ET_A$ ) by *Viburnum odoratissimum* and evaporation from containers with artificial foliage between early Mar. 2005 and Mar. 2006. Daily  $ET_A$  of live plant and evaporation from control containers were calculated based on changes in mass determined from weighing lysimeters. The  $ET_A$  line is representative of nine plant replicates, whereas the evaporation line is representative of three replicates. The least significant difference (LSD) line is the F-protected least significant difference between two means at  $\alpha = 0.05$ .

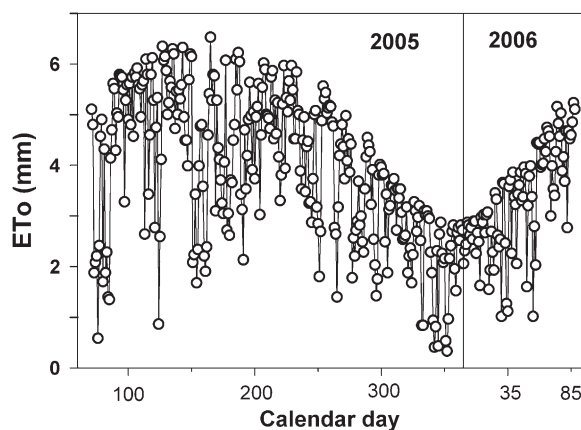


Fig. 2. Daily reference evapotranspiration ( $ETo$ ) from Mar. 2005 through Mar. 2006 at the Mid-Florida Research and Education Center, Apopka, FL. during the production of *Viburnum odoratissimum*.

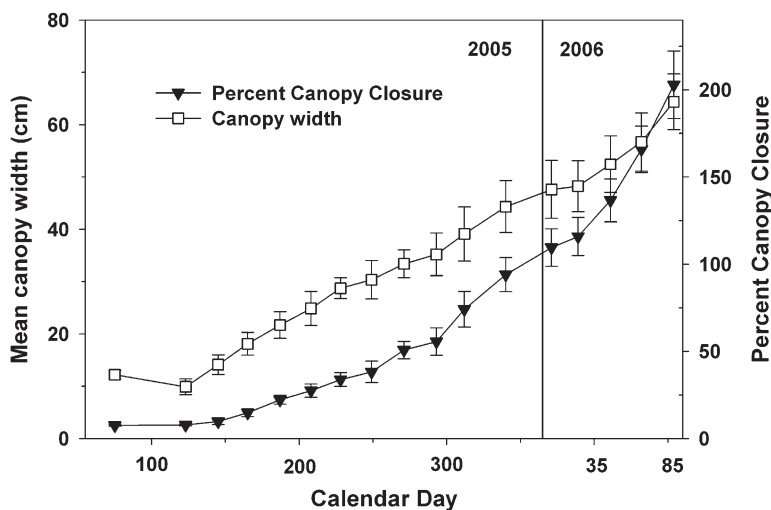


Fig. 3. Mean canopy widths measured for *Viburnum odoratissimum* and calculated %Closure from canopy widths and spacing between plants during production from rooted cuttings to market size plants for 13 months beginning in early Mar. 2005. Each mean is representative of nine single plant replicates. Error bars are the sd of nine plant replicates.



late Nov. 2005, mean %Closure was 94%. This level of %Closure likely also decoupled substrate evaporation, as shown previously for  $ET_A$ .

Plant transpiration dipped in Dec. 2005 as a result of shorter days, lower  $ET_o$  (Fig. 2), and termination of shoot growth in mid-December.  $ET_A$  increased in mid-February (Fig. 1) with increasing  $ET_o$  (Fig. 2) and initiation of bud break. This began a large and continuous shoot flush (Fig. 3), which doubled  $ET_A$  over the course of  $\approx 3$  weeks (Fig. 1). With this shoot flush, plants achieved market size by the end of March,  $\approx 13$  months after transplanting. As plants were approaching market size,  $ET_A$  was over a liter/day. In previous research (data not shown) with viburnum in the same containers with similar substrate and plant size, total plant-available water was determined to be 3.5 L (Beeson, 2007). Assuming similar plant-available water, at the peak in  $ET_A$  at market size, 30% to 40% of plant-available water was likely lost to evapotranspiration each day.

When summed over the production period, average  $ET_A$  was  $155 \pm 17.8$  L or  $3.08 \pm 0.35$  m in depth based on the top substrate surface area.  $ET_A$  ranged from 132.4 to 192.9 L per plant. Rainfall during production was 1.40 m. Although much of the rainfall supplemented overhead irrigation, rainfall in excess of container capacity was of no value. The capacity to retain rainfall would have varied day to day but would have generally increased as plants grew larger and transpired greater daily volumes of water. Nightly irrigating based on highest  $ET_A$  ensured that transpiration was fully coupled to  $ET_o$  for all plants. At harvest, canopy widths averaged 0.75 m and average height was 0.6 m. This was wider than the three-fourths height minimum (0.46 m) specified in the Florida Grades and Standards (DACS, 1994) and contributed to higher cumulative  $ET_A$  compared with  $142.8 \pm 9.1$  L ( $2.8 \pm 0.18$  m depth based on top substrate surface area) that was determined for *L. japonicum* in 2002 (Beeson, 2004).

The plot of WNI as a function of %Closure was fitted to an exponential decaying curve with an  $r^2 = 0.868$  and  $P = 0.001$  (Fig. 4). This curve was based on individual plants because PCA was not uniform. Thus, the equation models individual plant WNI rather than a population WNI, which accounts for some of the variability. WNI declined rapidly with small increases in plant size with the decline slowing  $\approx 20\%$  canopy closure at experimental spacing. Similar sharp declines for young plants were previously reported for *L. japonicum* (Beeson, 2004). For these viburnums, this slowing of the decline in WNI would have been when an average plant canopy covered  $\approx 80\%$  of the container substrate top surface, occurring around calendar Day 180.

The rapid decline in WNI at low canopy coverage occurred with increases in canopy growth for both species. Transpiration was strongly coupled to the environment characterized by  $ET_o$ , whereas substrate evaporation was not or was only weakly coupled as

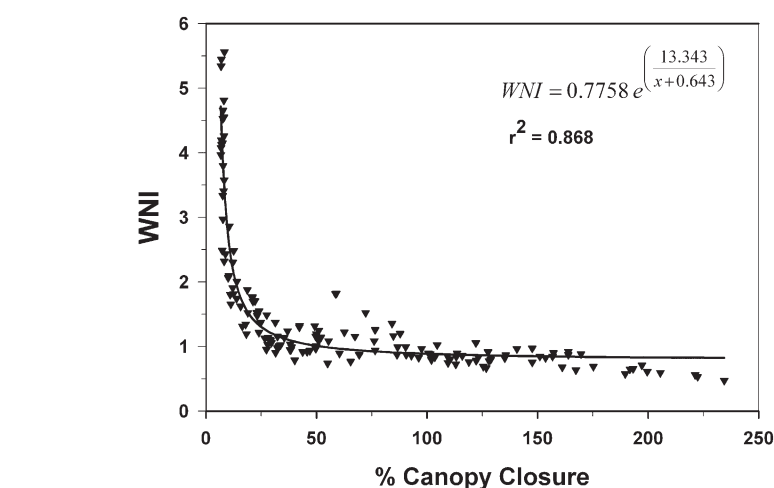


Fig. 4. Relationship between water needs index (WNI) values and corresponding %Closure for *Viburnum odoratissimum* during production from rooted cuttings to market size plants in 11.4-L containers. Points are the mean of individual weeks for each of nine individual plant replicates. The equation defines the decaying exponential curve that characterizes the relationship.

evidenced by consistent surface evaporation rates (Fig. 1). For young plants, PCA was quite small, 6% to 12% of the top substrate surface area (data not shown). The contribution of transpiration to  $ET_A$  was also quite small in contrast to surface evaporation. With  $ET_A$  driven by evaporation, not plant transpiration, but divided by a plant's small PCA compared with the much larger substrate surface area, calculated WNI was high. With plant growth, both PCA and transpiration increased, resulting in transpiration accounting for a larger portion of  $ET_A$ ; thus,  $ET_A$  became more responsive to  $ET_o$  resulting in lower WNI.

At  $\approx 70\%$  canopy closure, the rate of decline in WNI slowed as %Closure increased. This level of canopy closure is within the range where  $ET_A$  declines as a function of mutual sheltering among plants. At 100% canopy closure,  $ET_A$  was 40% lower for the same plants than at 67% closure or less (Beeson, 2010). Percent closures greater than 100% occur during production when plant canopies overlap. The greater the overlap, the denser and more uniform the overall canopy becomes, increasing overall boundary layer resistance to water vapor transfer (Noble, 1999). Increased resistance to vapor transfer lowers  $ET_A$  of individual plants, seen here as declines in WNI as %Closure increases. Nearly identical responses were observed for *L. japonicum* in the previous experiment (Beeson, 2004). For forest trees, Rose (1984) predicted this response based on shading of lower leaves and strong reductions in horizontal air movement through the lower canopy. Immediately after forests are thinned, the mirror image effect occurs as transpiration (Bréda et al., 1995) and boundary layer conductance of water vapor (Teklehaimanot et al., 1991) increase as a result of greater coupling of a tree's entire canopy to atmosphere rather than just the upper portion.

WNI calculated here and for *L. japonicum* is severalfold higher for young plants and

proceeds in the opposite direction of Kc calculated for agronomic crops (Allen et al., 1998) and other woody species in containers (Burger et al., 1987; Niu et al., 2006; Regan, 1997; Schuch and Burger, 1997). Crop coefficients increase with plant growth over time as a result of increased leaf area. Here, however, WNI is plotted against %Closure, not time. Although %Closure is related to time, it functions differently. Percent closure was chosen for several reasons. Foremost it is a direct function of plant growth and therefore independent of plant growth rate and regional climates. When plant growth arrests resulting from dormancy or episodic cycles, WNI does not change. Plant transpiration is assumed to remain coupled to the environment characterized by  $ET_o$  to the same degree with existing leaves exhibiting a consistent response. This eliminates need for Fourier curve transformations to account for changes in  $ET_A$  or growth when production extends across winter dormancy as proposed by Schuch and Burger (1997). Theoretically, %Closure should also be independent of container size and spacing within limits. For evergreen plants such as viburnum or *L. japonicum*, the %Closure model functions year-round, except when plants are exposed to sub-freezing temperature (Beeson, unpublished data). For deciduous shrubs, the %Closure model would only be applicable when plants are nearing or in full leaf. Factors that govern  $ET_A$  for evergreen plants would be similar to deciduous shrubs with a full canopy.

Data from the *L. japonicum* study were overlain with the viburnum results reported here to compare response of WNI to %Closure between species and years (Fig. 5). Statistics calculated for the combined data set by SigmaPlot (SSPS) indicate the coefficients for the resulting equation were significant ( $P < 0.0001$ ). The exponential decay equation for combined data had an  $r^2 = 0.843$ . This suggests that the combined equation could be used to determine WNI

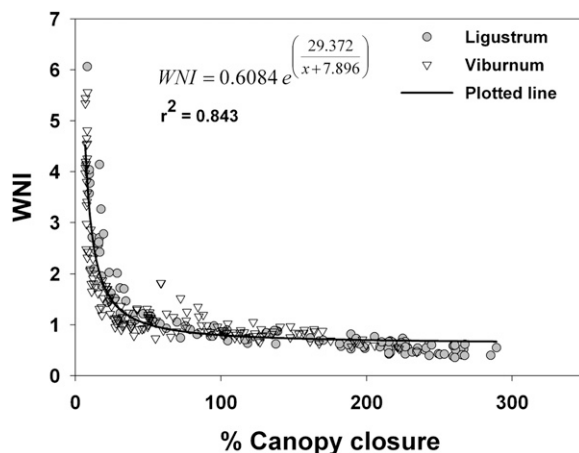


Fig. 5. Overlay of calculated water needs index (WNI) versus %Closure for *Ligustrum japonicum* data collected in 2001 to 2002 and *Viburnum odoratissimum* data collected in 2005 to 2006. Each point represents data from nine plant replicates over 7 continuous days. The equation was derived from combined data shown.

required for estimating irrigation requirements for either species. This also suggests that the %Closure model may be an avenue for the grouping of woody shrub species into general irrigation regimes based on canopy size and spacing.

The %Closure model presented here offers a way to relate ETo to plant size and spacing for calculation of precise irrigation rates from simple measurements. The process is based on plant canopy responses to the environment and lends itself to automation. Once the relationship between %Closure and WNI is established, estimation of daily ETo can be easily accomplished through spreadsheet calculations or with a computer algorithm. In practice, an irrigation manager would input container spacing likely once or twice during production, mean plant canopy width triweekly, with only ETo input daily. The greatest short-term variability in a crop's water use is directly related to the variability in solar radiation, which varies not only daily, but regionally. ETo in northwest Oregon can be 50% higher than in central Florida during the summer, mainly as a result of day length. Thus, actual ETo reported here is marginally consistent at 32° latitude and a coarse estimate 800 km north. Models based on physiological principles are the best avenue to make limited lysimeter data available to nursery industry worldwide.

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