Nitrogen Fertilization and Irrigation Frequency Affect Hydrangea Growth and Nutrient Uptake in Two Container Types

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Abstract. Plant growth, water use, photosynthetic performance, and nitrogen (N) uptake of ‘Merritt’s Supreme’ hydrangea (Hydrangea macrophylla) were investigated. Plants were fertilized with one of five N rates (0, 5, 10, 15, or 20 mM from NH₄NO₃), irrigated once or twice per day with the same total amount of water, and grown in either a paper biodegradable container or a traditional plastic container. Greater N rate generally increased plant growth index (PGI) in both plastic and biocontainers. Leaf and total plant dry weight (DW) increased with increasing N rate from 0 to 20 mM and stem and root DW were greatest when fertilized with 15 mM and 20 mM N. Plants fertilized with 20 mM N had the greatest leaf area and chlorophyll content in terms of SPAD reading. Container type had no influence on DW accumulation or leaf area. N concentrations (%) in leaves, roots, and the entire plant increased with increasing N rate. N concentrations in roots and in the entire plant were lower in biocontainers compared with plastic containers. Greater N rate generally increased daily water use (DWU), and biocontainers had greater DWU than entire plant were lower in biocontainers compared with plastic containers. Greater N rate and the entire plant increased with increasing N rate. High N fertilization rates (210 and 280 mg L⁻¹) increased plant N content and improved flowering performance, number of flowers, and flower size in ‘Merritt’s Supreme’ (Bi and Scagel, 2008); however, N leachate resulting from excessive N application can cause contamination of groundwater, which is not environmentally sustainable or cost efficient for growers. Concerns have been raised regarding N runoff from nursery production and possible environmental contamination (Yeager et al., 1993). An efficient fertilizer management program involves specific knowledge on the rate and application method of N fertilizer as well as the plant growth response to such fertilization practices.

Plant nutrient uptake is also affected by irrigation method (Scagel et al., 2011, 2012). Increased irrigation frequency was reported to increase N use efficiency and uptake of calcium (Ca) and decrease uptake of phosphorus (P), potassium (K), boron (B), and zinc (Zn) in Rhododendron species (Scagel et al., 2011, 2012). When the total amount of irrigation water was delivered through more than one irrigation event, it decreased leaching from containers and compensated for certain nutrient deficiencies (Scheiber et al., 2008; Silber et al., 2003; Xu et al., 2004). The effect of irrigation frequency on nutrient uptake was attributed to possible altered N availability in the substrate and to differences in plant biomass among treatments.

It is challenging for growers to determine water requirements of a specific species because it is not clear how much water is available for uptake in soilless substrates and how water status affects plant growth and nutrient uptake (O’Meara et al., 2014). Plant species vary in their ability to absorb water and nutrients as growing substrate dries out. O’Meara et al. (2014) assessed how decreasing substrate volumetric water content (VWC) influenced water uptake of H. macrophylla ‘Fasan’ (hydrangea) and Gardenia jasminoides ‘Radicans’ (gardenia). Water use by hydrangea started to decrease at a higher VWC (0.28 m³·m⁻³) than gardenia (0.20 m³·m⁻³) and water uptake ceased at 0.16 m³·m⁻³ in hydrangea, suggesting that hydrangea was less adept at extracting water from a drying substrate than the gardenia (O’Meara et al., 2014). For hydrangea, a relatively higher water content in the growing substrate should be maintained to maximize plant growth and avoid water stress. Sustainable alternatives have been studied in recent years to reduce use of plastic.
containers in nursery and greenhouse production of a number of ornamental crops (Beeks and Evans, 2013; Evans et al., 2010; Koeser et al., 2013; Kuehny et al., 2011; Nambuchiri et al., 2015; Wang et al., 2015). Biodegradable containers, also known as biocontainers, are made from a variety of biodegradable materials, such as feather, fabric, rice hulls, and paper, thus introducing varying influence on plant growth and nutrient uptake. Many tested biocontainers are able to produce plants of similar quality to traditional plastic containers (Beeks and Evans, 2013; Koeser et al., 2013; Kuehny et al., 2011; Li et al., 2015). However, compared with plastic containers, some biocontainers require more frequent irrigations and increased amounts of irrigation water when they are made from porous hydrophilic materials, for instance paper, which has high evaporation loss through the container sidewall (Evans et al., 2010; Koeser et al., 2013; Wang et al., 2015). The increased water use in biodegradable containers, mostly considered to be an insignificant contribution to total production cost (Brumfield et al., 2015), has an unforeseen impact on water status and nutrient availability in the substrate and, therefore, plant growth and nutrient uptake.

The objectives of this study were as follows: 1) to investigate plant growth and nutrient uptake of ‘Merritt’s Supreme’ hydrangea in response to N fertilization rate, irrigation frequency, and container type; and 2) investigate water use of hydrangea growing in black plastic containers compared with biodegradable containers made from recycled paper.

Materials and Methods

Plant culture and treatments. One hundred rooted liners of ‘Merritt’s Supreme’ hydrangea were transplanted into two types of containers on 1 July 2014: a black plastic container (GL 400; top diameter 17.78 cm, bottom diameter 18.10 cm, volume 3.785 L; Nursery Supplies Inc., Chambersburg, PA), or a biodegradable container (also referred as biocontainer) made from a mix of recycled paper (7 x 7 round; interior top diameter 18.7 cm, bottom diameter 14.9 cm, height 17.1 cm, volume 3.90 L; Western Pulp Products Co., Corvallis, OR). Hydrangea plants were maintained outdoors under a shade structure with 50% black shadecloth at the R.R. Foil Plant Science Research center of Mississippi State University (U.S. Department of Agriculture hardiness zone 8a; 33.4552° N, 88.7944° W). A substrate containing ≈60% composted pine bark, 30% sphagnum peatmoss, and 10% perlite by volume (Metro Mix 852; Sun Gro Horticulture, Agawam, MA) was used as the growing substrate. Each hydrangea plant was fertilized with 250 mL N-free fertilizer (1.06 mg·mL⁻¹; Cornell No. N Eq. 0–6–27; GreenCare Fertilizers, Kan-kakee, IL) (Table 1) twice weekly plus 0.5, 10, 15, or 20 mM N from NH₄NO₃ from 8 July to 22 Sept. 2014. Plants were irrigated through drip irrigation either once per day at 0800 hr or twice per day at 0800 and 1430 hr with the same total daily irrigation volume. Plants were irrigated to replace daily water loss plus 10% to 15% leaching. Water loss was determined gravimetrically, and irrigation volume was corrected according to the average of the plant height and two widths. Leaf chlorophyll content was estimated by leaf SPAD reading. Leaf SPAD for an individual plant was measured from three fully expanded new leaves every 2 weeks using a chlorophyll meter (SPAD 502 Plus; Konica Minolta, Osaka, Japan). Three readings from the three selected leaves were averaged to represent leaf SPAD of a specific plant.

Photosynthetic measurements. Leaf net photosynthetic rate and gs of hydrangea plants were measured between 1000 and 1300 hr on 27 Aug., 11 Sept., 22 Sept., and 8 Oct. 2014 using a portable photosynthesis system (LI-6400XT; LI-COR Biosciences, Lincoln, NE). One of the first two pairs of fully expanded leaves, not shaded by other leaves, were selected for photosynthetic measurements on a specific plant (Currey and Lopez, 2015). The selected leaf was enclosed into a 2-cm² leaf chamber with a fluorometer light source. Photosynthetically active radiation of 1000 µmol·m⁻²·s⁻¹ and a reference CO₂ concentration of 400 µmol·mol⁻¹ were maintained inside the leaf chamber during photosynthetic measurements. Block temperature inside the leaf chamber was maintained according to the atmosphere temperature on the measurement date.

Daily water use and substrate moisture. Daily water use was measured in plants irrigated once per day. Daily water use was determined using a gravimetric method by subtracting pot weight (plant included) 24 h after irrigation from pot weight at container capacity (approximately one-half hour after irrigation). Substrate moisture at 5-cm depth in each container was measured using a soil moisture sensor (WaterScout SM 100; Spectrum Technologies, Inc., Aurora, IL). The moisture sensor was connected to a soil sensor reader (FieldScout® 6466; Spectrum Technologies, Inc.) for instant substrate moisture readings. Substrate moisture was measured at the midpoint between the plant stem and container sidewall after the second weighing of the container, 24 h after the first irrigation following DWU measurement. Daily water use and substrate moisture were measured on four dates: 21 Aug., 9 Sept., 25 Sept., and 9 Oct. 2014.

Plant harvest. Each plant was destructively harvested on 27 Oct. 2014. 119 d after transplanting. Plant samples were cleaned free of any substrate with deionized water, and separated into leaf, stem, and root tissues. All leaves from one plant were passed through a leaf area meter (LI-3100C; LI-COR Biosciences) to measure the total leaf area of each plant. Roots of three freshly harvested plants from each treatment combination were scanned (EPSON® Expression 10000XL; Epson America, Inc., Long Beach, CA) for an image. Root length and surface area were analyzed using root analysis software (WinRHIZO, Regent Instruments Inc., Québec, QC, Canada). All samples were oven dried at 60 °C. Dry weight of each sample was recorded. Total DW of a specific plant was calculated by summing the DW of leaf, stem, and root tissues.

Tissue nitrogen analyses. Each dry sample was ground to pass a 1-mm sieve (Wiley Mill; Thomas Scientific, Swedesboro, NJ) for nutrient analyses. The kjeldahl method was used for determination of total N concentration (%) using 0.1 g of dry tissue (Bremner, 1965). Nitrogen content (mg) was calculated by multiplying DW of each sample with its N concentration. Total N content of each plant was calculated by summing N content in leaf, stem, and root tissues. Average N concentration of a plant was then determined by dividing total plant N content by total plant DW.

Experimental design and data analyses. This study was set up as a factorial arrangement of treatments in a completely randomized design. The N rate (five rates), container type (two types), and irrigation frequency (two frequencies) were the three main experimental factors providing 20 treatment combinations with five single-plant replications in each treatment combination. Significance of any main effect or the interaction among factors were determined by analysis of variance (ANOVA) using the GLM procedure of SAS (Version 9.4; SAS Institute, Cary, NC). Daily water use and substrate moisture data on different dates were subject to repeated measures with measurement date as a factor. Where indicated by ANOVA, means were separated by Fisher’s protected least significant difference test at P < 0.05.

Results

Plant growth index. The influence of N rate on PGI varied between container type on most measuring dates (Fig. 1; PGI shown only for 19 Aug. and 27 Oct.). In each container type, PGI increased with increasing N rate from 0 to 15 mM N on 19 Aug. and 27 Oct. On 19 Aug., after 42 d of treatment, biocontainers resulted in 18.5%, 9.8%, and 7.6% greater PGI than plastic containers at
10, 15, and 20 mM N, respectively. By the end of the experiment on 27 Oct., PGI was similar between container types at any N rate from 5 to 20 mM. Irrigation frequency influenced PGI only at 42 d (19 Aug.) when two irrigations per day produced plants with 5.9% higher PGI than one irrigation per day (Fig. 1C).

Leaf SPAD. Relative leaf chlorophyll content (estimated via SPAD) was affected by N rate on all measurement dates (Fig. 2A; SPAD shown only for 27 Oct. at the end of the season). Plastic containers increased SPAD readings by 7.0%, 4.4%, 3.4%, and 4.4% compared with biocontainers on 24 July, 21 Aug., 5 Sept., and 3 Oct., respectively (data not shown). Leaf SPAD was not influenced by irrigation frequency. On 27 Oct., plants fertilized with 20 mM N had the greatest SPAD readings. Leaf SPAD readings between plants fertilized with 5, 10, or 15 mM N were similar, all of which resulted in higher SPAD readings than the no N treatment.

Dry weight and leaf area. Greater N rate increased total plant DW and leaf DW from 0 to 20 mM N (Fig. 3). Stem and root DW increased with increasing N rate from 0 to 15 mM, with no difference between plants fertilized with 15 or 20 mM N. Leaf area of hydrangea plants also increased with increasing N rate from 0 to 20 mM N (Fig. 2B). Container type and irrigation frequency had no influence on plant DW or leaf area.

Root length and surface area. The influence of N rate on total root length varied between container types (Fig. 4A). In general, 10 to 20 mM N resulted in the greatest root lengths and 0 mM N resulted in the least in both plastic and biocontainers. Biocontainers resulted in greater root length than plastic containers at 5 mM N. Root length peaked at 15 mM N in biocontainers and at 10 mM N in plastic containers, resulting in similar root length at N rates of 10 to 20 mM in biocontainers or plastic containers. Irrigation frequency altered the effect of N rate on total root length (Fig. 4B). Nitrogen rates of 15 and 20 mM resulted in greater root length than 0 or 5 mM N regardless of irrigation frequency. Irrigation frequency influenced root length only at 10 and 15 mM N. More frequent irrigation increased root length at 10 mM N and had the opposite effect at 15 mM N. Root surface area was affected by N rate, but not by container type or irrigation frequency (Fig. 4C). Root surface area increased with increasing N rate from 0 to 15 mM N, with no difference between plants fertilized with 15 or 20 mM N.

Daily water use and substrate moisture. Daily water use increased with N rate from 0 to 15 mM N, with 15 and 20 mM N resulting in similar DWU (Fig. 5A). Greater DWU was consistent with increased PGI and dry weight of hydrangea plants using similar high N rates. Biocontainers resulted in 16.2% greater DWU than plastic containers (Fig. 5B). In general, substrate moisture decreased with increasing N rate, with no difference between 5 and 10 mM N, or between 15 and 20 mM N (Fig. 6A). Nitrogen rates of 15 mM and 20 mM resulted in the lowest substrate moisture. Plastic containers had 10.1% lower substrate moisture at 5-cm depth than biocontainers (Fig. 6B). More frequent irrigation resulted in 13.2% greater substrate moisture before onset of scheduled irrigation in the morning (Fig. 6C).

Photosynthetic rate. The influence of N rate on net photosynthetic rate varied between container types on 27 Aug. (Fig. 7A). Plastic containers resulted the highest net photosynthetic rate on 27 Aug. at N rates of 10 to 20 mM, higher than that of plants grown in biocontainers at similar N rates as well as those fertilized with 0 or 5 mM N grown in plastic or biocontainers. Nitrogen rate affected net photosynthetic rate measured on 11 Sept., 22 Sept., and 8 Oct. (Fig. 7B–D). On 11 Sept. and 22 Sept., N rate of 20 mM resulted in higher net photosynthetic rate than 0 and 5 mM N, with no N resulting in the lowest net photosynthetic rate. On 8 Oct., net photosynthetic rates were generally higher when plants were fertilized with N than the no N treatment. In addition, on 8 Oct., there was a container effect on net photosynthetic rate where net photosynthetic rate was 10% greater in plastic containers than in biocontainers (Fig. 7E).

Irrigation frequency did not affect net photosynthetic rate of hydrangea plants in this study on any measurement date.

Stomatal conductance. The effects of N rate on gs varied among the four measuring dates (Fig. 8). On 27 Aug., plants fertilized with N from 5 to 20 mM had similar gs, and gs of N-fertilized plants was higher than those fertilized with no N. On September measurement dates, gs generally increased with N rate (11 Sept.) or had little influence on gs (22 Sept.). At the end of the study, N rates from 0 to 10 mM resulted in higher gs than the rates of 15 or 20 mM N on 8 Oct. Plants grown in plastic containers had greater gs than plants in biocontainers on the three dates measured. Irrigation frequency did not influence gs.

Tissue N concentration. Generally, increasing N rate resulted in increasing N concentration in leaf and root tissues, as well as the plant average, with 20 mM N producing
the highest N concentrations in leaf, root, and the plant average (Fig. 9). Stem N concentration was unaffected by N rate. One irrigation per day increased stem N concentration by 6.4% compared with two irrigations per day. Plastic containers resulted in 10.3% and 9.6% higher root and average plant N concentrations than biocontainers, respectively.

Discussion

Hydrangeas are generally considered to have high nutrient requirements, especially N, to support their vigorous growth (Bi and Scagel, 2008; Bi et al., 2008). Greater N rate increased plant biomass, leaf quality, tissue N content, and flower number and size in ‘Merritt’s Supreme’ hydrangea (Bi et al., 2008), which is consistent with our results, in which higher N rates produced plants with greater PGI, leaf area, dry weights, and tissue N concentrations. Similar results between N rate and growth are also reported for azalea (Rhododendron sp.) and ‘Bartlett’ pear (Pyrus communis) (Bi et al., 2007; Cheng et al., 2001). High tissue N content in late
summer or fall is considered beneficial for spring growth and flowering the following season that is supported by the remobilization of stored N (Bi et al., 2003; Cheng and Xia, 2004; Millard, 1995). On the other hand, runoff of N to the environment due to over-application of fertilizer can cause potential contamination of groundwater and should be avoided. According to our results, 20 mM N resulted in the greatest plant DW, leaf SPAD reading, leaf area, and tissue N concentration (in leaf, root, and average plant). Plant growth index, root length, and surface area were comparable with 15 and 20 mM N in plastic or in biocontainers. These results suggest that the optimal N rate for 'Merritt's Supreme' may be greater than the highest rate used in our study, but will depend on the variables used to assess plant quality. Because our study was focused on vegetative growth period, decisions on the optimum N rate should include flowering performance in the following season and whether nursery growers are marketing flowering plants or not.

In this study, more frequent irrigation increased substrate moisture but had little influence on plant growth or nutrient uptake. Scagel et al. (2012) reported that the effect of irrigation frequency on nutrient uptake in rhododendron (Rhododendron sp.) may be a result of effect on plant biomass, rather than a result of substrate nutrient availability or plant uptake ability. The accumulation of N in plant tissue is regulated by the growth rate and rate of biomass accumulation of a given species, especially under sufficient N supply (Gastal and Lemaire, 2002). In our study, two irrigations per day were designed to decrease leaching of nutrients to the environment and alleviate possible plant water stress during hot summer conditions by maintaining a higher substrate moisture level. Although two irrigations per day increased substrate moisture at 5-cm depth, gs, an indicator of plant water status, was unaffected by irrigation frequency, suggesting increased substrate moisture with two irrigations per day may not have improved water status (or reduced water stress) of hydrangea plants.

Substrate moisture at 5-cm depth generally decreased with increasing N rate, with 15 and 20 mM N resulting in the lowest moisture. Plant water uptake through the root is mainly

Fig. 6. Substrate moisture of Hydrangea macrophylla ‘Merritt’s Supreme’ affected by N rate (A), container type (B), or irrigation frequency (C). Hydrangea plants were fertilized with 0, 5, 10, 15, or 20 mM N from NH4NO3, grown in plastic container or paper biocontainer, and irrigated once or twice per day with the same total daily irrigation volume. Substrate moisture was measured at 5-cm depth using a soil moisture sensor. Different lowercase letters in a figure suggest significant difference compared by Fisher’s least significant difference test at \( P < 0.05 \).

Fig. 7. Net photosynthetic rate of Hydrangea macrophylla ‘Merritt’s Supreme’ affected by the interaction between N rate and container type on 27 Aug. (A), by the main effect of N rate on 11 Sept. (B) and 22 Sept. (C), or by the main effects of N rate (D) and container type (E) on 8 Oct. with no interactions between main effects. Hydrangea plants were fertilized with 0, 5, 10, 15, or 20 mM N from NH4NO3, grown in plastic container or paper biocontainer, and irrigated once or twice per day with the same total daily irrigation volume. Different lowercase letters in a figure suggest significant difference among all treatment combinations compared by Fisher’s least significant difference test at \( P < 0.05 \).
Fig. 8. The stomatal conductance ($g_s$) of *Hydrangea macrophylla* ‘Merritt’s Supreme’ affected by N rate (A–D), or by container type (E–G) on different measurement dates. Hydrangea plants were fertilized with 0, 5, 10, 15, or 20 mM N from NH$_4$NO$_3$, grown in plastic container or paper biocontainer, and irrigated once or twice per day with the same total daily irrigation volume. Irrigation frequency did not affect $g_s$. Different lowercase letters in a figure suggest significant difference among N rates or between container types compared by Fisher’s least significant difference test at $P < 0.05$.

Fig. 9. Tissue N concentration of *Hydrangea macrophylla* ‘Merritt’s Supreme’ affected by N rate in the plant, leaves, and roots (A–D), by container type in the plant (E) and roots (G), or by irrigation frequency in stems (F). Hydrangea plants were fertilized with 0, 5, 10, 15, or 20 mM N from NH$_4$NO$_3$, grown in plastic container or paper biocontainer, and irrigated once or twice per day with the same total daily irrigation volume. Different lowercase letters in a figure suggest significant difference among N rates, between container types, or between irrigation frequencies compared by Fisher’s least significant difference test at $P < 0.05$. 

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lost through transpiration and evaporation (Jason and Lawlor, 1979; McElrone et al., 2013). Hydrangea plants fertilized with 15 and 20 mM N had higher PGI and possibly transpire more water than plants fertilized with N rates of 0 to 10 mM N. Therefore, lower substrate moisture with 15 and 20 mM N was consistent with high transpiration and water use of these plants. Two irrigations per day increased substrate moisture. With one irrigation per day, lower moisture at 15 and 20 mM N did not limit plant root growth, possibly because plants were established to have a robust root system. Two irrigations per day increased root length at 10 mM N compared with one irrigation. It is possible that increased moisture level promoted plant root growth when it was not as established as at 15 or 20 mM N, considering that 15 and 20 mM N resulted in higher root DW than 10 mM N. However, altered substrate moisture did not affect N uptake measured by tissue N concentration, except for increased stem N concentration with two irrigations per day.

The paper biocontainer used in this study increased DWU of hydrangea plants compared with plastic containers. Wang et al. (2015) also reported increased DWU in plants grown in wood pulp containers compared with plastic containers. The wood pulp container used by Wang et al. (2015) is made from similar materials and manufactured by the same company as the paper biocontainers used in our study. However, substrate moisture at 5-cm depth was higher in biocontainers than in plastic containers. In a traditional plastic container, evaporation is mainly through the substrate surface. Evaporation occurred through both substrate surface and container sidewalls in the paper biocontainers. This characteristic may have contributed to increased DWU besides plant transpiration. Higher substrate moisture at 5-cm depth in biocontainers also may result from the considerable evaporation through the container sidewall, leaving high moisture level toward the substrate surface. By comparison, moisture is mainly lost through the substrate surface in plastic containers, which resulted in the lower moisture at 5-cm depth. Therefore, the different evaporation patterns in the two container types may have caused the higher substrate moisture at 5-cm depth in paper biocontainers than plastic containers. Nambuthiri et al. (2015) found lower substrate temperature and lower sidewall temperature in a wood pulp alternative container than in black plastic containers. The contribution of sidewall water loss to overall container evapotranspiration has a major influence on reducing substrate temperature and the evaporative cooling effect, associated with increased water use, and may help reduce heat stress and enhance plant survival in locations with high summer temperatures (Nambuthiri et al., 2015). When grown in anebb-and-flood subirrigation system, ‘Rainier Purple’ cyclamen (Cyclamen persicum) had higher dry root weights when grown in paper and wood fiber containers than in plastic containers (Beeks and Evans, 2013). Beneficial effects on root growth were also found on ‘Chiffon’ Encore® azalea when grown in paper biocontainers compared with black plastic containers (Li et al., 2018). In the current study, container type had little influence on plant growth. Even though biocontainers had greater substrate moisture at 5 cm than plastic containers, biocontainers resulted in a higher DWU and lower N uptake. These results indicate that although both container types produce similar plants after 3 months, using biocontainers will require a greater amount of water and may have a higher potential for nutrient leaching from the substrate.

The paper biocontainers resulted in lower gs and much lower net photosynthetic rate than in plastic containers, indicating that the increased water use in biocontainers caused greater water stress and decreased opening of stomata in hydrangea plants. Rose et al. (1990) reported plant growth of some woody ornamentals, such as crabapple (Malus × zumi) and maple (Acer × freemontii E.), were enhanced more by minimizing water stress than by increasing fertilizer concentration. Biocontainer-produced hydrangea plants were of comparable visual quality as plastic container–produced plants, but had decreased N concentration in the root and in the plant on average. With ‘Chiffon’ Encore® azalea, plants in paper biocontainers had increased PGI, plant dry weight, and plant N content at 10, 15, or 20 mM N rates, possibly due to azalea’s preference for good drainage and aeration, which is provided by the paper biocontainers (Li et al., 2018). However, biocontainers decreased N uptake in hydrangea plants in our study. It is possible the increased water stress in biocontainers limited N availability or the root’s ability to uptake N. Leaf SPAD has been used to help predict leaf N status in a number of species (Heerema et al., 2014; Jifon et al., 2005; Kim et al., 2013; Netto et al., 2005). Lower leaf SPAD readings in hydrangea plants grown in biocontainers may be a reflection of the reduced N uptake. Irrigation frequency did not affect plant photosynthetic rate, gs, or N uptake in the entire plant. Although two irrigations per day resulted in higher substrate moisture than one irrigation per day, such effect did not reduce the water stress caused by biocontainers.

Optimal environment for photosynthesis varies among plant species (Salisbury and Ross, 1992). Day temperature affects photosynthesis, where net photosynthetic rate increases with increasing temperature and starts to drop when reaching a critical high temperature (Armitage et al., 1990; Lasseigne et al., 2007). Under fluctuating environmental conditions, photosynthesis of hydrangea plants was affected by container type, N rate, or the interaction between N rate and container type rather than by irrigation frequency. Photosynthetic capacity was reported to have a strong correlation with plant N status, especially in leaves (Evans, 1989). Heerema et al. (2014) showed a decrease in photosynthesis in pecan (Carya illinoensis) leaves with decreasing leaf N when N was remobilized from leaves for fruit development. Nitrogen rate in our study affected net photosynthetic rate and gs of hydrangea plants on all measuring dates. The highest net photosynthetic rate was found at 20 mM N on 11 Sept. and 22 Sept., suggesting sufficient N fertilization is needed to support photosynthesis in hydrangea plants. Decreased gs was reported in response to a short-time water stress with ‘Better Boy’ tomato plants (Lycopersicon esculentum), in which the depressing effect of water stress on gs was reversed after rewatering (Gu et al., 1996). Growing hydrangea in plastic containers generally increased net photosynthetic rate and gs. Lower gs of plants in paper biocontainers indicated greater water stress. Such container effect is possibly derived from increased DWU in biocontainers and decreased N uptake by the hydrangea plants. Although the effects of container type on DWU, substrate moisture, and N uptake did not result in major differences in plant growth after 3 months, these differences might influence future plant growth and quality in hydrangea plants.

In conclusion, ‘Merritt’s Supreme’ hydrangea has a high N requirement where 20 mM N resulted in the greatest plant dry weight, leaf SPAD readings, leaf area, and N concentration (in leaf, root, and the entire plant). N rate of 15 mM resulted in similar PGI, root length, and surface area, and dry weights of stem and root as 20 mM N. The paper biocontainers used in our study increased DWU while reducing plant photosynthetic rate, gs, N uptake, and leaf SPAD readings in hydrangea plants compared with plastic containers. Two irrigations per day resulted in greater substrate moisture at 5-cm depth than one irrigation, but increased irrigation frequency did not compensate for the increased water loss from biocontainers because irrigation frequency did not affect net photosynthetic rate or gs. Considering the negative effects of water use and nutrient uptake of plants grown in the paper biocontainers, use of traditional plastic containers is recommended for nursery production of ‘Merritt’s Supreme’ hydrangea.

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