Impact of Applied Nitrogen Concentration on Growth of Elatior Begonia and New Guinea Impatiens and Susceptibility of Begonia to Botrytis cinerea

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ABSTRACT. Plant performance and appearance in deficient and toxic levels of nutrients are well characterized. However, less is known about the potential subtleties of plant growth, form, development, nutrient uptake, and biotic stress tolerance in the broad tolerable range. Begonia [Begonia × tuberhybrida Voss] and new guinea impatiens [NGI (Impatiens hawkeri Bull.)] were grown over a wide range of N (from 1.78 to 57.1 mM NH₄NO₃ ratio at a 1:1 ratio supplied as nutrient solution) in a peat/perlite soilless substrate in greenhouse conditions. Plant growth, development, chlorophyll content, leaf angle, nutrient uptake, tissue caloric value, and susceptibility to Botrytis cinerea Pers.:Fr. disease were evaluated in two experiments. Elevated N supply resulted in decreased plant height (16% in Beg and 7% to 16% in NGI), flower count (3% to 48% in Beg and 7% to 49% in NGI), bud numbers (23% to 80% in Beg), canopy area (11% to 33% in NGI), and mass (21% to 33% in Beg and 18% to 58% in NGI). Chlorophyll content saturated at an N supply of 28.6 mM. N uptake efficiency, shoot N use efficiency, and shoot N utilization efficiency decreased with increasing N supply. Elevated levels of N supply from 7.15 to 57.1 mM also increased the susceptibility of Beg to B. cinerea disease by 10% to 80% in stems and 3% to 14% in leaves. The increase in susceptibility also corresponded with increased tissue energy content (kJ g⁻¹) and altered leaf orientation. This study indicates many plant changes occur between nutrient extremes that can have a significant impact on growth, development, and the ability to withstand disease.

Nitrogen uptake by plants is generally in anionic (NO₃⁻), cationic (NH₄⁺), or neutral (CH₃N₂O) forms. Excessive addition of a single element may cause an imbalance of other nutrients (Marshner, 1995; Mengel and Kirkby, 2001; Mills and Jones, 1996) and may predispose plants to disease (Engelhard, 1989; Hoffland et al., 2000; Huber and Watson, 1974; Jarvis, 1992; Mansfield, 1980; Marshner, 1995). The probability of N toxicity is rare because unlike micronutrients, a toxic N concentration is several-hundred-fold higher than sufficient N concentrations. However, the practice of periodic fertigation (fertilizer mixed with irrigation water) of concentrations substantially higher than published recommendations is not uncommon under intensive greenhouse production systems. The N use efficiency is critical in determining N supply rate, which ultimately can reduce input costs while protecting the environment from N deposition (Good et al., 2004). In greenhouse production systems, especially under soilless culture, plant N uptake is solely dependant on external supply. Objective assessment of N use efficiency will provide vital information for the greenhouse industry, which leads to prudent management of resources.

Much of the literature on plant nutrition focuses on plant growth effects in the nutritional extremes of either toxicity or deficiency. There are fewer evaluations of the sometimes subtle changes in plant growth and appearance and the potential for increased plant pathogen susceptibility in commonly encountered N concentrations (Kent and Reed, 1996; Nelson et al., 1978; Smith et al., 1998). Studies have linked N supply and disease (David et al., 2003; Engelhard, 1989; Huber and Watson, 1974; Mengel and Kirby, 2001). High N supply has resulted in increased disease susceptibility to Pseudomonas syringae van Hall pv. tomato (Okabe) Young, Dye & Wilkie and Oidium lycopersicum Cook & Mass. for tomato (Lycopersicum lycopersicum L.) (Hoffland

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et al., 2000) in contrast with resistance to *B. cinerea* with high N supply in tomato (Hoffland et al., 1999). This inconsistency could be the result of the type of pathogens, N form, source and concentration, type of substrate, irrigation frequency, and growing environment. Commercially blended fertilizers that contain high concentrations of NH$_4^+$ are lower in Ca (Nelson, 1996). High levels of Ca and K have significant influence on disease resistance (Elad and Volpin, 1993; Marshner, 1995; Mengel and Kirby, 2001). Calcium has a major physiological role in forming the cell membrane and cell wall, which protects against pathogens (Elad, 1988; Elad and Evensen, 1995; Volpin and Elad, 1991). High potassium also suppresses disease in plants (Mengel and Kirby, 2001; Talbot and Zeiger, 1996). These confounding findings could be a potential factor for conflicting reports of the influence of N in disease-related studies.

*B. cinerea* is a ubiquitous pathogen that infects leaves, stems, and flowers. Symptoms of the disease appear as water-soaked lesion spots, which quickly coalesce affecting the entire tissue, causing major economic loss (Elad, 1988; Hausbeck and Moorman, 1996). Susceptibility to *B. cinerea* disease has been reported to decrease with NO$_3^-$ form but increase with NH$_4^+$ form (Huber and Watson, 1974). Conversely, it has also been reported that *B. cinerea* shows no growth benefit based on the form of N supplied to the plant (Townsend, 1957).

Begonia [Begonia × tuberhybrida Voss] and new guinea inpatiens [NGI (*Impatiens hawkeri* Bull.)] are categorized as high-valued plants grown in pots and hanging baskets (U.S. Dept. Agr., 2005). These plants are succulent with tender tissues and are considered inefficient in terms of water use, requiring ample water supply but not water-logged conditions (Hartley, 1995). This creates a moist growing environment causing the plants to be susceptible to the ubiquitous fungal pathogen *B. cinerea* (Elad and Shtienberg, 1995; Hausbeck and Moorman, 1996). The nutritional requirement is moderate for these species (Nelson, 2005). All these factors contributed to the selection of these greenhouse plants as model species for biotic stress and nutritional interaction studies.

This study was conducted to investigate the growth of Beg and NGI, N uptake efficiency, shoot N use efficiency, and shoot N utilization efficiency and the potential susceptibility of these plants to biotic stress (disease: *B. cinerea*) to N concentrations that are commonly found in commercial greenhouses. This study provides insight about the nutritional economy of plants across a range of N supply.

**Materials and Methods**

Two experiments were conducted in a glass greenhouse set at night/cloudy day/clear day temperatures of 17/21/24 °C with relative humidity ranging from 20% to 50%. In Expt. 1, NGI cv. Pure Beauty Purple was grown from 22 Jan. 2004 to 4 Apr. 2004, and in Expt. 2, Beg cv. Barbara was grown from 7 Apr. 2004 to 1 June 2004. Photoperiod was maintained at 16 h using a 1 lamp:1 lamp ratio of high-pressure sodium and metal halide lamps. Minimum light levels were 200 μmol·m$^-2$·s$^-1$ (under electric lamps) and as high as 800 μmol·m$^-2$·s$^-1$ during sunny days measured with a single quantum sensor (model QSO; Apogee Instruments, Logan, Utah) located in the center of the greenhouse. In both experiments, rooted cuttings were transplanted into 15-cm diameter pots containing 70% sphagnum peatmoss, 30% horticultural grade perlite (by volume) amended with dolomitic limestone (3.0 mg·L$^-1$), and a micronutrient mix (0.59 mg·L$^-1$, 12% Fe, 2.5% Mn, 1.0% Zn, 0.5% Zn, 0.1% B, and 0.05% Mo by mass, Micromax; Scott’s Co., Marysville, Ohio). The newly transplanted cuttings were watered with reverse osmosis water once and then fertilized with 3.6 mM nutrient concentration of 20N–8.8P–16.6K fertilizer (0.5% Mg, 0.0068% B, 0.0036% Cu, 0.05% Fe, 0.025% Mn, 0.0009% Mo, 0.0025% Zn; Scott’s Co.) during the initial 2 weeks of establishment before treatments. Treatment nutrient solution pHs were adjusted to 5.8 using Ca(OH)$_2$ and H$_2$SO$_4$ as required. The plants were irrigated to runoff and received at least a 15% leachate fraction (estimated) to reduce the risk of continuous salt buildup resulting from the fertilizer treatments. Plants were harvested during the seventh and eighth weeks of treatment for Beg and NGI, respectively. Some of the Beg plants were set aside at that time to study their susceptibility to *B. cinerea*.

In Expt. 1, the treatments consisted of six N levels, 1.78, 3.57, 7.15, 14.3, 28.6, and 42.86 mm N as NH$_4$NO$_3$ in the nutrient solution. Expt. 2 consisted of the same N treatments plus 57.1 mm N in the same form. The remainder of the nutrient solutions consisted of millimolar concentrations of 0.67 PO$_4$-P, 1.67 K, 2.3 Ca, 1.3 Mg, and 1.3 SO$_4$-S plus micromolar concentrations of 71 Fe, 9 Mn, 1.5 Cu, 1.5 Zn, 45 B, 0.1 Mo, 4.6 Cl, and 0.2 Na using KH$_2$PO$_4$, MgSO$_4$, K-EDTA, CaCl$_2$, Fe-DTPA, MnCl$_2$, CuCl$_2$, ZnCl$_2$, H$_2$BO$_3$, and Na$_2$-MoO$_4$. The solutions used deionized water of 18 Mohm purity. The mean electrical conductivity (EC) of the treatment nutrient solutions with 1.78, 3.57, 7.15, 14.3, 28.6, 42.86, and 57.1 mm N were 1.5, 1.8, 2.2, 2.8, 3.3, 4.3, and 4.8 mS·cm$^-1$, respectively. The experiments were laid out in randomized complete block designs with six (Expt. 1) or seven (Expt. 2) treatments and five replications with three plants per experimental replicate.

**Growth measurements.** In both experiments, the plants were photographed and analyzed for canopy coverage using image analysis software (Assess; American Phytopathological Society, St. Paul, Minn.). Percent maximum canopy coverage of leaf canopy area was calculated by dividing the total number of pixels of each treatment by the highest pixel treatment. Leaf orientation angle was determined by superimposing a protractor, in software, to a plant’s side view and measuring three to four mature or maturing leaves from three replicate plants in each treatment. At the end of the growth period, plant height was measured from the base of the stem to the top of the plant, flower and bud number were counted, and then harvested by cutting the stem of the plant at the top of the soilless medium. Shoots were dried in a forced air drying oven at 70 °C for at least 2 d and dry weight was recorded. There were no apparent differences in the starting size of the plants for height, mass, spread, and leaf angles.

**pH and electrical conductivity measurements.** Substrate pH and EC values were measured using the pour-through method (Wright, 1986), whereby 50 mL of leachate was collected from each pot by adding 150 mL of deionized water on the surface of the substrate; an additional 20 mL to 30 mL of water was added if there was insufficient leachate as a result of dryness of the substrate. The pH and EC values were measured from the collected leachate using electric probes (models Inlab 730 and 413; Mettler Toledo, Columbus, Ohio) and a meter (model SevenMulti; Mettler Toledo).
**Chlorophyll measurement.** Chlorophyll content was estimated nondestructively using a portable chlorophyll meter (SPAD-502; Minolta Camera Co., Ltd., Tokyo). Four readings per plant were taken on the adaxial surface of recently matured leaves.

**Energy measurement.** Caloric values were measured according to the method described by Long (1934) and Yang et al. (2003). Oven-dried ground tissue samples of 1 g or less were used. A known amount of mineral oil was added to the sample as a binding agent and pressed into a pellet with a press (Parr Instrument Co., Moline, Ill.). The pellets were then combusted in a bomb calorimeter (Parr 6200; Parr Instrument Co.). Because mineral oil burns with the plant sample, the effect of mineral oil combustion must be corrected by multiplying mineral oil energy content by mineral oil amount in each sample and subtracting that value from the total energetic value. The difference is the energetic content of the sample. The total shoot caloric value of a plant was calculated by multiplying the caloric value per gram of tissue by total shoot dry weight.

**Tissue analysis.** Recently matured leaf samples were washed in 0.1 N HCl for 30 s and rinsed in deionized water for 30 s. The samples were oven-dried at 70 °C for at least 2 d and ground in a mortar and pestle. The tissue was analyzed for macro- and micronutrients with the exception of N using inductively coupled plasma emission spectrophotometry (model IRIS Intrepid II; Thermo Electron Corp., Waltham, Mass.). Total N was analyzed using a C-H-N analyzer (Carlo Erba NA 1500 Nitrogen Analyzer; Thermo Electron Corp.).

*Botrytis cinerea* inoculation. *Botrytis cinerea* was cultured on potato dextrose agar plates (DIFCO, Sparks, Md.) at 25 °C for 7 d to 10 d until it produced abundant conidia. The plates were flooded with reverse osmosis water and conidia dislodged with a glass rod. The conidial suspension was adjusted to $2.5 \times 10^6$ conidia/mL using a hemacytometer and sprayed onto plants to runoff. Beg plants were inoculated during the eighth week at the time other plants were harvested and measured for growth. Figure 1 shows the approximate plant sizes of each treatment. The inoculated plants were kept in a plastic-covered tunnel that maintained at least 80% relative humidity using a humidifier. The inoculated plants were incubated for 10 d. On day 10 after inoculation, the plants were scored for black to gray necrotic lesions on the leaf lamina and stems.

**Nitrogen uptake efficiency.** Each efficiency parameter was defined previously by Goins et al. (2004) and Good et al. (2004). N uptake efficiency (tissue N concentration × shoot dry weight/N supply) is a measure of total N taken up and accumulated by the plant in relation to the total N supplied. Shoot N use efficiency (shoot dry weight/N supply) measures the change in shoot dry weight per unit of N supplied to the plant. Shoot N utilization efficiency (shoot dry weight/total N content) × 100] measures the efficiency of N partitioning to the shoot.

**Statistical analysis.** All the data were subjected to analysis of variance using PROC GLM SAS (SAS Institute, Cary, N.C.). Where the F test indicated a significant ($P \leq 0.05$) treatment effect, differences among the treatment means were computed by LSMEANS and compared by the least significant difference 0.05 method. Regression analysis was done to assess...
linear and quadratic effects using Sigma Plot (version 6.0; Jandel Scientific, San Rafael, Calif.).

**Results**

**Plant Growth.** Plant height peaked at 7.15 mm N and decreased at higher N rates for Beg (Figs. 1A, 2). The reduction in plant height was more gradual and linear in NGI with a height difference of 13.6% between the lowest and the highest N supply. In Beg, the highest N treatment resulted in shoots that were comparable in size to the lowest N treatment but ≈3 cm or 15% shorter than the 7.15 mm N treatment. The number of flowers or buds was highest in both species at low N supply (1.78 to 7.15 mm N; Fig. 1B) and declined (3% to 48% in Beg and 7% to 49% in NGI) as N supply increased above 14.3 mm.

There was also a peak in shoot dry weight at 7.15 mm N (Fig. 1C), but additional N supply resulted in decreased shoot dry weight (21% to 33% in Beg and 18% to 58% in NGI). The lowest shoot dry weight was produced at highest N level of 42.86 and 57.1 mm in NGI and Beg, respectively. The shoot dry weight produced at the lowest N was greater than at the highest N for both species.

Leaf orientation also changed with different N supply rates. Leaf orientation was more upright at the lowest N concentration and less upright as the N concentration increased (Figs. 1D and E, 3A–D). The leaf angle decreased as N supply increased within different leaf age groups in both species. Leaf orientation was almost horizontal, less than 20° and 10° for maturing leaves and matured leaves, respectively. Leaf angle was greater in young and maturing leaves for both crops with N levels less than 14.7 mm. This was best seen in Beg plants by looking at the uppermost (young, expanding) leaves. When viewed from above the plant, many leaf edges can be seen in plants receiving low N concentrations, whereas in high N concentrations, full leaves arranged parallel to the growing surface could be seen (Fig. 3A and B). For NGI, the leaf angle is best seen from the side as leaves of all ages were droopy and curled when fed with increasing N concentrations (Fig. 2).

Chlorophyll content, based on SPAD readings, increased for both Beg and NGI with higher N (Table 1). Chlorophyll content reached a maximum at 28.6 and 42.86 mm N for NGI and Beg, respectively. NGI leaf canopy area was greatest at 7.15 mm N and decreased significantly at higher N (Table 1). Maturing and recently matured leaves of NGI developed necrosis resembling K deficiency or NH₄⁺ toxicity along the margins from the...
Table 1. Correlation between N supply and chlorophyll content as indicated by SPAD value and percent maximum canopy coverage in begonia (Beg) and new guinea impatiens (NGI).

<table>
<thead>
<tr>
<th>Treatment</th>
<th>SPAD value</th>
<th>Maximum canopy coverage (%)</th>
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<tbody>
<tr>
<td>N supplied (mm)</td>
<td>Beg</td>
<td>NGI</td>
</tr>
<tr>
<td>1.78</td>
<td>42.6</td>
<td>62.5</td>
</tr>
<tr>
<td>3.57</td>
<td>49.5</td>
<td>69.5</td>
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<tr>
<td>7.15</td>
<td>53.7</td>
<td>70.4</td>
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<tr>
<td>14.3</td>
<td>56.9</td>
<td>76.3</td>
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<td>28.6</td>
<td>59.2</td>
<td>78.3</td>
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<tr>
<td>42.86</td>
<td>62.6</td>
<td>76.9</td>
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<td>57.10</td>
<td>63.0</td>
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| P | ** | * | * |
| Linear | ** | * | * |
| Quadratic | NS | * | NS |

$R^2 = 0.8547, 0.9097, 0.8859$ *Nonsignificant or significant at $P < 0.05$ or 0.01, respectively.

terminal tip to the middle of the leaf lamina with N 28 mm or greater N. High N (greater than 28 mm N) plants had poor root systems. Beg plants treated with high N developed black necrotic lesions at the stem base close to the substrate. Maturings and matured leaves of Beg treated with 1.78 mm N developed light chlorosis with reddish pigmented margins resembling visual symptoms of N deficiency (Sprague, 1964). These symptoms were not apparent in NGI, perhaps as a result of the use of a dark-leaved variety in this study.

**Botrytis cinerea: Begonia Interaction.** When Beg plants were inoculated with *B. cinerea*, disease severity increased. The low N supply treatments from 1.78 to 7.15 mm N had less than 5% infection on leaves and stems (Fig. 1F). Plants fertigated with 28.6 mm N and above had higher percentages of infection, especially on the stems. Stem lesions occurred at the basal portion of the plant, which caused the plants to collapse. In some instances, plant death followed rapid *B. cinerea* progression.

**Nutrient Interactions.** There was a negative quadratic correlation between pH and increasing N supply from the third to seventh weeks for both species (Fig. 4A). The fertilizer used in this study would decrease the pH if NH$_4^+$ was preferentially taken up by the plant over NO$_3^-$. If this was the case, the resulting acidity would quickly neutralize the lime added to the media in our study. As a result of the nature of the fertilizer treatments, the EC also increased with higher N supply (Fig. 4B), which can also cause pH to drop. The EC of the fertilizer collected from pots was consistently lower for NGI than Beg. This difference may have been attributable in part to differences in nutrient uptake between the species. For each macronutrient, NGI contained more than Beg (Fig. 5A–E and results below), which would lead to less salts collected from the NGI pots.

Tissue N concentration increased quadratically with increasing N supply in both species (Fig. 5A). In NGI tissue, N concentrations were 2.6% and 7.5% at N concentrations of 1.78 and 42.86 mm, respectively. In Beg, the tissue N concentrations were 2.0% and 5.5% with N concentrations of 1.78 and 57.1 mm, respectively. In NGI, tissue P concentration increased quadratically with increasing N supply. Tissue Mg peaked at the highest value at 14.1 mm N (Fig. 5B) but decreased slightly at higher N concentrations. Tissue P increased linearly with increasing N for Beg. There was a significant linear decrease in K content with increased N supply in Beg (Fig. 5C). NGI tissue Ca was highest between 7.15 and 14.3 mm N in NGI (Fig. 5D) and then declined. Beg tissue Ca increased quadratically with increasing N supply. Tissue Mg peaked at 7.15 mm N for both species (Fig. 5E). The shapes of the curves could be a function of many different conditions in the root zone. Saturation of uptake may be occurring with some nutrients; nutrient availability could reach equilibrium in midranges of N concentrations (along with the corresponding EC and pH changes), or antagonistic/protagonistic effects may be altered as N concentration and preference for different N forms changes.

Significant differences in tissue Fe, Mn, Zn, Cu, and B concentration were noted with increasing N supply for both species (Fig. 6A and B). With some elements, the statistical trend is hidden as a result of the scale on the y-axis, so the regression line, $r^2$, and $P$ value are shown for clarity.

**Tissue Energy Value.** Beg tissue energy content increased from 15.1 kJ·g$^{-1}$ (1.78 mm N) to 19.7 kJ·g$^{-1}$ (57.1 mm N). Total shoot energetic value peaked at 7.15 mm N [242 kJ·plant$^{-1}$ (Beg)] and dropped to 167 kJ (Beg) at 57.1 mm N (NGI) (Fig. 7B). No such statistically significant trend was measured for the NGI.

Fig. 4. pH and electrical conductivity (EC) of collected leachate from pots of the different N treatments of begonia (Beg) and new guinea impatiens (NGI). Regression models between N supply and pH values (A) in Beg n = 29; $y = 6.23 - 0.18x + 0.0029x^2$ and NGI n = 29; $y = 6.49 - 0.14x + 0.0035x^2$, or EC values (B) in Beg n = 29; $y = 0.93 - 0.016x + 0.0015x^2$ and NGI n = 29; $y = 1.62 + 0.092x - 0.0002x^2$. $r^2$ values are presented when it is statistically significant at $P < 0.05$. Error bars are ±1 standard deviation from the average.

Fig. 5. pH and electrical conductivity (EC) of collected leachate from pots of the different N treatments of begonia (Beg) and new guinea impatiens (NGI). Regression models between N supply and pH values (A) in Beg n = 29; $y = 6.23 - 0.18x + 0.0029x^2$ and NGI n = 29; $y = 6.49 - 0.14x + 0.0035x^2$, or EC values (B) in Beg n = 29; $y = 0.93 - 0.016x + 0.0015x^2$ and NGI n = 29; $y = 1.62 + 0.092x - 0.0002x^2$. $r^2$ values are presented when it is statistically significant at $P < 0.05$. Error bars are ±1 standard deviation from the average.

Fig. 6. Electrical conductivity (EC) of collected leachate from pots of the different N treatments of begonia (Beg) and new guinea impatiens (NGI). Regression models between N supply and pH values (A) in Beg n = 29; $y = 6.23 - 0.18x + 0.0029x^2$ and NGI n = 29; $y = 6.49 - 0.14x + 0.0035x^2$, or EC values (B) in Beg n = 29; $y = 0.93 - 0.016x + 0.0015x^2$ and NGI n = 29; $y = 1.62 + 0.092x - 0.0002x^2$. $r^2$ values are presented when it is statistically significant at $P < 0.05$. Error bars are ±1 standard deviation from the average.

Fig. 7. Electrical conductivity (EC) of collected leachate from pots of the different N treatments of begonia (Beg) and new guinea impatiens (NGI). Regression models between N supply and pH values (A) in Beg n = 29; $y = 6.23 - 0.18x + 0.0029x^2$ and NGI n = 29; $y = 6.49 - 0.14x + 0.0035x^2$, or EC values (B) in Beg n = 29; $y = 0.93 - 0.016x + 0.0015x^2$ and NGI n = 29; $y = 1.62 + 0.092x - 0.0002x^2$. $r^2$ values are presented when it is statistically significant at $P < 0.05$. Error bars are ±1 standard deviation from the average.
NITROGEN UPTAKE, USE, AND UTILIZATION EFFICIENCY. The N uptake, use, and utilization efficiency decreased with increasing N for both species (Fig. 8A–C). N uptake was most efficient at 1.78 mM N, but at 14.3 mM N, shoot use efficiency was close to zero in both species. The shoot N use efficiency was lower than the N uptake efficiency by over 50% in lower N supply treatments. The shoot N use efficiency was greater in Beg than NGI from 1.78 to 7.15 mM N supply. At 14.3 mM N supply, shoot N use efficiency reached close to zero in both species. Shoot N utilization efficiency reached its lowest value of ≈20% for Beg and 15% for NGI at or above 28.6 mM N supply.

Discussion

Plants supplied with N rates greater than 14.3 mM N were more compact, had darker green leaves, and fewer flower buds and flowers (Fig. 1) than plants at lower N rates indicating that and NGI (Fig. 4), and this may have resulted in additional negative effects on nutrient uptake (Epstein and Bloom, 2005; Marshner, 1995) and growth.

Nitrogen is the only element that exists in the environment in cationic and anionic forms, and N uptake will alter the rhizosphere pH potentially resulting in synergistic or antagonistic effect on uptake of other cations and anions (Hamlin et al., 1999; Kirby and Mengel, 1967; Magalhaes and Huber, 1989). NH$_4^+$ supply may have suppressed K, Ca, and Mg uptake (Fig. 5), which resulted in low tissue concentrations of the respective elements in both species, although experimental levels are within critical guidelines for these species (Mills and Jones, 1996). However P, an anion, was above normal tissue P concentration of 0.52% (Pitchay, 2002). The synergistic effect on P uptake was noted at N supply of 28.7 msior less. However, the high EC and low buffering capacity of the substrate (Fig. 4; Harris et al., 1999; Osorio et al., 2003) may have resulted in decreased availability and uptake of competing nutrients.
High rates of N increased the number of plants infected with *B. cinerea* with high mortality occurring above 28.7 mM N (Fig. 1). Hoffland et al. (2000) reported similar findings in tomato, and David et al. (2003) reported the same pattern in powdery mildew (*Erysiphe cichoracearum* DC) infection in Beg. Several factors could have contributed to the increase in infection with increasing N supply treatments. The reduction in the shoot dry weight at supraoptimal N levels may have altered the C:N ratio at the cellular level. Free amino acids and low-molecular-weight nitrogenous compounds are good energy sources for pathogens (Hoffland et al., 1999). The preferential uptake of NH$_4^+$ over NO$_3^-$ and antagonistic effect on K, Ca, and Mg uptake may have caused significant ionic imbalances affecting cell homeostasis (Britto and Kronzucker, 2005; Kirby and Mengel, 1967). The relatively low tissue concentration of K and Ca and the unbalanced N:K ratio in the nutrient solution as a result of increased N supply could have weakened the cell membrane and affected cell wall development (Stall, 1963; Volpin and Elad, 1991) resulting in less resistance to pathogen penetration of the cell walls. Damaged membranes could have also resulted in nutrient leakage that provided stimulation of *B. cinerea* infection (Harrison, 1988). The increase in tissue energy values of Beg plants supplied with elevated N (Fig. 7), indicating a nutrient-rich environment, increases the likelihood of successful colonization resulting from larger energy supplies per gram of biomass consumed by the pathogen. On the other hand, the low energy value of low N-supplied plants may have been an unfavorable environment for *B. cinerea* infection.

The change in the leaf angle and orientation with different levels of N supply, especially the leaf phylloplane (Fig. 1), may have provided a conducive environment for the pathogenic spores to land and attach themselves as an initial step of establishment (van Kan, 2005). This is followed by the anchoring and growing of the conidia as it uses the nutrients deposited on the leaves, which was more likely to be abundant in the horizontal phylloplane (as a result of residual deposition without runoff from leaves from overhead fertigation) as compared with more upright leaves.

Poor root systems in plants supplied with high N could have affected water uptake resulting in the substrate remaining moist for prolonged periods. This contributed to higher humidity microenvironments in these plants, especially near the lower portion of the plants, which are ideal for disease infection (Harrison, 1988).
N uptake efficiency decreased with increasing N supply (Fig. 8). In a supraoptimal N environment, the additional C needed for amino acid synthesis was likely provided at the expense of biomass synthesis (Champigny, 1995). This shift in carbon partitioning may result in N within the tissue not being assimilated because the N uptake mechanism remains efficient, whereas dry matter gain is less efficient. The linear increase in energy per gram of tissues from plants supplied with elevated N rate (Fig. 7) may have contributed to the corresponding high infection rate of *B. cinerea* in Beg. 

N use efficiency declined significantly with increasing N supply (Fig. 8). Higher levels of N increased tissue caloric value and altered leaf orientation, which correlated with greater disease incidence. N supply between the ranges of 3.57 and 14.3 mM was found to be optimal for plant growth and development in Beg and NGI based on appearance, growth, and development and susceptibility to *B. cinerea* infection. These amounts differ from amounts commonly applied in commercial greenhouses of at least 14.3 mM with periodic fertigation double or triple that amount. Additional studies on N form and concentration and interaction of environmental factors with species that are sensitive to both extremes of N form will generate more information for N use efficiency. Additional tests should be done to determine the direct versus indirect effects of elevated N on floricultural crops to see if elevated N primarily caused the growth effects and increased susceptibility or the resulting stresses from altered pH and EC caused the observed effects. Regardless of the specific causes, elevated yet sublethal N levels can lead to significant growth effects on these floricultural crops.

**Literature Cited**


