Response of *Oxypetalum caeruleum* to Irradiance, Temperature, and Photoperiod

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Abstract. Incremental increases in temperature from 14 to 22 to 30°C resulted in linear increases in stem length and node number and decreases in stem diameter and stem strength of *Oxypetalum caeruleum* (D. Don.) Decne. Higher temperatures also resulted in additional flower abortion, reduced time to flowering, and fewer flowering stems per inflorescence. Reduction in the photosynthetic photon flux (PPF) from 695 to 315 µmol·s⁻¹·m⁻² had similar effects as increasing the temperature on vegetative characteristics, but had little effect on reproductive ones. The rate of stem elongation was greatest at low PPF for all temperatures and at high temperature for all PPF treatments. Net photosynthesis rose between 14 and 22°C and declined at 30°C for all PPF treatments. Long photoperiods (12 or 14 hours) resulted in longer internodes, longer stems, and more flowers per cyme than short photoperiods (8 or 10 hours), but photoperiod had little effect on flowering time. Treatments to reduce latex coagulant and silver thiosulfate treatments had no significant effect on vase life.

*Oxypetalum caeruleum* (tweedia) belongs to Asclepiadaceae and bears light-green opposite leaves and cymose panicles, each cyme consisting of three to five five-petaled light-blue flowers. The species is native to Brazil and Uruguay (Bailey, 1951) and grown in the United States, Holland, New Zealand, and Australia as a specialty cut flower crop. Other important ornamental crops in Asclepiadaceae are *Asclepias tuberosa L.*, *Stephanotis floribunda* Brongn., and *Hoya carnosa* R. Br. *Asclepias* is day-neutral with respect to flowering, but long days promote shoot extension and leaf production (Lyns, 1985) and inhibit lateral branching (Lyns and Booze, 1983). *Stephanotis* is a long-day plant for flower initiation, with an optimum of 21°C for flowering (Kofranek and Criley, 1983), and *Hoya* has a minimum of 15°C for flowering (Post, 1955). Little literature is available on the response of *Oxypetalum* to light and temperature, although Post (1955) suggested a minimum of 18°C for best vegetative growth. Similarly, no studies have been conducted to determine photosynthetic characteristics of the species.

Cut stems of *Oxypetalum* exude a milky latex-like sap. Halsey and Mayak (1981) noted that many flowers with milky latex tended to have short vase lives, which was generally attributed to latex plugging of the conducting vessels. Isopropyl alcohol (Freyermuth et al., 1984; Gordon et al., 1986), a boiling water dip (Salinger, 1985; Stirling, 1950; Systems, 1966), or a warm water dip (45°C) for 15 min (Salinger, 1985) has been proposed to eliminate the exudate. Vase life of numerous species has been extended with silver thiosulfate (STS) (Farnham et al., 1981; Veen and van de Geijen, 1978); proprietary preservative solutions, mainly those containing sucrose and biocides, are also commonly employed. The objectives of this work were to 1) demonstrate the effects of photosynthetic photon flux (PPF) on flowering, quality of flowering stems, and photosynthesis of tweedia, 2) demonstrate the effects of temperature and photoperiod on growth and flowering, and 3) evaluate the vase life of the cut flowers and determine if vase life could be extended by selected conditioning treatments.

Materials and Methods

Light-temperature studies. Rooted cuttings were received from Gunderson Nurseries, Otaki, New Zealand, and all stems were immediately pinched to two nodes to ensure removal of reproductive tissue. Seventy-two uniform plants growing in 1.2-liter-capacity pots containing 5 peat : 5 pumice (v/v) growing medium were randomly placed in nine light–temperature combinations. Plants were grown in walk-in controlled-environment (CE) rooms at the Dept. of Scientific and Industrial Research Climate Laboratory, Palmerston North, New Zealand. Temperatures were constant at 30, 22, or 14 ± 0.5°C. A constant vapor pressure deficit of 0.45 kPa was maintained resulting in relative humidities of 90%, 83%, or 72% ± 5%, respectively. Lighting over the 12-hr photoperiod was provided by four 1000-W high-pressure multivapor lamps (Sylvania “Metalarc”) and four 1000-W Philips quartz halogen lamps separated from the chamber by a plate glass and water thermal barrier. The PPF at pot surface height was 695 ± 20 µmol·s⁻¹·m⁻² (high light). Use of woven Saran cloth of =35% density provided a PPF of 450 ± 10 µmol·s⁻¹·m⁻² (medium light), and cloth of 55% density, a PPF of 315 ± 8 µmol·s⁻¹·m⁻² (low light). Each density of Saran cloth was suspended over one PPF treatment in each of the growth rooms. Additional information on room design, temperature inputs, humidity control, and lighting design has been described previously (DSIR Climate Laboratory, 1981; Barrington and Kanemasu, 1983). Plants were watered with 100 ml of complete mineral nutrient solution (half-strength Hoagland’s A) (Hoagland and Arnon, 1938) three times daily from an automated irrigation system.

Stem length and number of nodes were recorded once a week on four randomly selected plants at the beginning of the exper-

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results from the main treatments are presented. The reduction in temperature from 30 to 14 °C resulted in a linear decrease in stem length and number of nodes and a linear increase in stem diameter and stem strength (Table 1). Stem length was significantly affected by temperature; stems grew 38% longer at 30 °C than at 14 °C. The additional stem length resulted from additional nodes and longer internodes. Average internode length was 4.7 cm at 30 °C and 4.5 cm at 14 °C. The significant quadratic trend for node number was due to the small difference between 14 and 22 °C, compared with the large difference between 22 and 30 °C. Critical load (stem strength) is a function of stem length and stem diameter (Goeschl et al., 1966). The decrease in stem length and coincident increase in stem diameter with decreasing temperature resulted in a linear increase in critical load (Table 1).

Reducing PPF produced the opposite effect of decreasing temperature. The decrease in PPF from 695 to 315 µmol·s⁻¹·m⁻² resulted in a linear increase in stem length and node number and a linear decrease in stem diameter, number of stems, and stem strength (Table 1). Stem length was also affected by PPF; each 10-µmol·s⁻¹·m⁻² reduction in PPF resulted in a 0.5-cm increase in stem length due to an increase in node number and internode length. Average internode lengths were 4.1 and 5.2 cm for PPF values of 695 and 315 µmol·s⁻¹·m⁻², respectively. The number of stems was significantly reduced as PPF decreased but was unaffected by temperature (Table 1), indicating that PPF was more important in stem initiation and development than temperature.

Rate of stem elongation. Stems elongated sigmoidally over time (Fig. 1, upper and lower left). Stem elongation was more rapid the higher the temperature and was most rapid between weeks 1 and 4 for plants grown at 30 and 22 °C, regardless of PPF. Plants grown at 14 °C, however, did not elongate rapidly until after week 4, but sustained that rate for an additional 7 to 10 weeks, depending on PPF treatment. The rates of stem elongation during the main phase of growth were determined by fitting linear regressions of stem length over time during the period 0.25 to 0.75 of the interval from the beginning of treatments to flowering (McPherson et al., 1985). Stem elongation rate at 14 °C was significantly lower, regardless of PPF treatment, compared with stems grown at 22 or 30 °C (Fig. 1, lower right). Stems grew slowest on plants under high PPF and fastest under low PPF for all temperatures (Fig. 1, lower right). The increase

Table 1. Effect of temperature and PPF on flowering stem characteristics of Oxypetalum caeruleum. No interactions occurred for any characteristic. The values were recorded at time of first flowering in each treatment.

| Variable       | Stem length (cm) | Stem diam (cm) | Stems (no.) | Nodes per stem (no.) | Critical load (mm·cm⁻²) |
|----------------|------------------|----------------|-------------|----------------------|------------------------|
| Temperature    |                  |                |             |                      |                        |
| 30             | 62.2             | 29.7           | 2.6         | 13.3                 | 0.48                   |
| 22             | 50.4             | 30.9           | 2.6         | 10.5                 | 0.61                   |
| 14             | 45.1             | 35.4           | 2.5         | 10.1                 | 0.78                   |
| T¹linear       | *                | *              | NS          | *                    | *                      |
| T²quadratic    | NS               | NS             | NS          | *                    | NS                     |
| PPF (µmol·s⁻¹·m⁻²) |      |                |             |                      |                        |
| 695            | 43.4             | 33.5           | 3.9         | 10.7                 | 0.77                   |
| 450            | 54.3             | 32.0           | 2.1         | 11.3                 | 0.59                   |
| 315            | 62.4             | 30.3           | 1.5         | 12.0                 | 0.48                   |
| T¹linear       | *                | *              | *           | *                    | *                      |
| T²quadratic    | NS               | NS             | NS          | NS                   | NS                     |

Results and Discussion

Light × temperature

Vegetative characteristics. As there were no significant temperature × PPF interactions for any vegetative criterion measured, only the results from the main treatments are presented. The reduction in temperature from 30 to 14 °C resulted in a linear decrease in stem length and number of nodes and a linear increase in stem diameter and stem strength (Table 1). Stem length was significantly affected by temperature; stems grew 38% longer at 30 °C than at 14 °C. The additional stem length resulted from additional nodes and longer internodes. Average internode length was 4.7 cm at 30 °C and 4.5 cm at 14 °C. The significant quadratic trend for node number was due to the small difference between 14 and 22 °C, compared with the large difference between 22 and 30 °C. Critical load (stem strength) is a function of stem length and stem diameter (Goeschl et al., 1966). The decrease in stem length and coincident increase in stem diameter with decreasing temperature resulted in a linear increase in critical load (Table 1).

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in stem elongation between 14 and 30°C was lower for high PPF (64%) than for medium (276%) and low PPF (178%).

Net photosynthesis. Pn differed significantly for all PPF treatments at 30 and 22°C; however, no differences occurred in Pn at 14°C, regardless of PPF (Fig. 2). Pn rose between 14 and 22°C and declined between 22 and 30°C for all PPF treatments (Fig. 2), although an optimum rate may have occurred between 24 and 26°C, particularly under high PPF conditions. This response to temperature is similar to that of other C₃ plants, where the optimum temperature for photosynthesis is in the 20 to 25°C range (Berry and Bjorkman, 1980). A high rate of photosynthesis is commonly observed in leaves acclimated to high, in contrast to low, PPF conditions (Boardman, 1977). Pn was 80% saturated at 1045 µmol·s⁻¹·m⁻² (Fig. 3), and the apparent photon quantum yield was =0.020 mol CO₂/mol photons. The saturation point was high compared with geraniums (Pelargonium x hortorum Bailey) and impatiens (Impatiens hybrida) (Armitage and Vines, 1982), but the apparent photon yield was low compared with other dicotyledonous C₃ species that have a photon yield (adsorbed light basis) of 0.052 mol CO₂/mol photons (Ehleringer and Pearcy, 1983).

Reproductive characteristics. No temperature x PPF interactions occurred for any reproductive characteristic measured. With the exception of aborted flowers, all criteria measured were far more responsive to changes in temperature than to changes in PPF (Table 2). Time to flowering was delayed as temperature was lowered, especially between 22 and 14°C (Table 2), indicating that Oxypetalum should not be grown at cool temperatures. Other genera of Asclepiadaceae show similar temperature effects on flowering. Stephanotis has an optimum of
Fig. 3. Effect of PPF on Pn of two single-plant replications of Oxypetalum caendeum grown at 22°C and PPF of 695 µmol·s⁻¹·m⁻². Each data point is the mean of two readings. + = 30°C, closed circles = 22°C.

Table 2. Effect of temperature and PPF on reproductive characteristics of Oxypetalum caeruleum. No interactions occurred for any characteristic. The values were recorded at time of first flowering in each treatment.

| Temperature (°C) | Time to flower (days) | Flowers per cyme | Aborted flowers | Flower nodes to produce six cymes |
|------------------|-----------------------|------------------|-----------------|----------------------------------|
|                  |                       |                  | (%), 3 to 5     | (%)                             |
|                  |                       |                  | (%), 1 to 2     | (%)                             |
|                  |                       |                  | (%), 6 cymes    | (%)                             |
| 30               | 32                    | 11.4             | 69.0            | 20.1                            | 8                              |
| 22               | 38                    | 70.0             | 26.0            | 3.9                             | 6                              |
| 14               | 115                   | 84.6             | 5.8             | 9.6                             | 6                              |
|                  |                       |                  |                 |                                 |                                |
|                  |                       |                  |                 |                                 |                                |
| PPF (µmol·s⁻¹·m⁻²) |                       |                  |                 |                                 |                                |
| 695              | 63                    | 64.0             | 30.5            | 6.4                             | 6                              |
| 450              | 59                    | 54.4             | 34.9            | 10.7                            | 6                              |
| 315              | 63                    | 47.2             | 34.3            | 18.4                            | 7                              |
|                  |                       |                  |                 |                                 |                                |
|                  |                       |                  |                 |                                 |                                |

Table 3. Influence of photoperiod on growth and flowering of Oxypetalum caeruleum.

| Photoperiod (hr) | Stems (no.) | Internode length (cm) | Time to flower (days) | Flowers per cyme |
|------------------|-------------|-----------------------|-----------------------|-----------------|
|                  | 3 to 5 (%)  | 1 to 2 (%)            |                       |                 |
| 8                | 39          | 58                    | 33                    |                 |
| 10               | 39          | 61                    | 38                    |                 |
| 12               | 48          | 48                    | 35                    |                 |
| 14               | 71          | 27                    | 41                    |                 |

Fig. 4. Influence of photoperiod on stem length and internode length of Oxypetalum caeruleum. (left to right) 8-, 10-, 12-, 14-hr photoperiod.

21°C (Kofranek and Criley, 1983) and Hoya has a minimum of 15°C for flowering (Post, 1955).

The morphology of the inflorescence was significantly affected by changes in temperature. In Oxypetalum, a well-developed inflorescence consists of cymes with four to five flowers. At low temperature (14°C), most cymes consisted of three to five flowers, few had one to two flowers, and few aborted. However, abortion was greater and the proportion of cymes with only one to two flowers were highest at 30°C (Table 2). Eight floral nodes were necessary on the panicle to form six cymes on plants grown at 30°C. The reduction in cymes as inflorescences increased in length at 30°C resulted in poorly developed flower stems and reduced ornamental value. The quadratic trends shown in the temperature data resulted from the relatively large differences between 30°C and 22°C, compared with the differences between 22 and 14°C for all criteria measured (Table 2). These data indicate that 14°C is too cold for commercial flowering and that 30°C results in high rates of flower abortion and poorly developed inflorescences. Temperatures close to 22°C appear optimum for flowering time and flower development. An additional benefit of 22°C was the effect on flower color. At 30°C, flowers were blue-pink and tended to fade rapidly, at 22°C flowers were sky blue, and at 14°C they were a muddy, less-attractive blue-mauve.

A trend toward a reduced number of cymes with three to five flowers and an increased number of aborted flowers per cyme occurred under low-PPF conditions (Table 2). In general, however, PPF affected vegetative characteristics more than reproductive parameters (Tables 1 and 2).

Photoperiod

Most characteristics measured showed a significant linear response to lengthening photoperiod (Table 3). The number of nodes were unaffected by photoperiod (data not shown) and the increase in stem length may be largely attributed to the increase in internode length (Table 3). The longest stems occurred under the longest photoperiods (Fig. 4). The number of flowers per cyme also increased with increasing daylength (Table 3). Stems with more flowers per inflorescence appeared fuller and were of better quality under longer photoperiods. In general, photo-
periods of 12 and 14 hr resulted in longer stems and higher flower quality than 8 and 10 hr, important attributes for cut flower crops. Stem diameter, stem strength, and aborted flowers were not significantly affected due to photoperiod treatments (data not shown).

Postharvest evaluations

Control stems persisted 8 days, a duration adequate for commercial purposes, but no latex coagulant treatment resulted in significant improvement in vase life. The average vase life for all treatments was 9 days. The use of boiling water did not enhance vase life; although, it was beneficial in handling stems due to the reduction of stickiness of the latex. Isopropyl alcohol caused rapid leaf abscission on some stems, but the damage was inconsistent. This result contrasts with those of Freyermuth et al. (1984), who found beneficial effects of isopropyl alcohol on poinsettia. No differences in vase life occurred due to STS treatments; STS-treated stems (4 or 8 mM) abscised slightly later, but flowers still reverted to pink at about the same time as those of control stems.

In summary, both temperature and PPF played a significant role in the growth and flowering of Oxypetalum caendeum. Temperature, however, affected flowering responses and flower morphology more than PPF. Photoperiod had little effect on flowering time but had a significant influence on internode elongation and stem morphology. Growth at 22°C and a 12-hr photoperiod resulted in excellent flower color, realistic flowering time, high critical load, and highest quality cymes. Postharvest experiments showed that latex from cut stems was not detrimental to the marketable life of the cut stem and treatments that reduced latex did not result in additional vase life.

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