Modeling the Effects of Temperature and Photosynthetic Daily Light Integral on Growth and Flowering of *Salvia splendens* and *Tagetes patula*

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**Abstract.** Photosynthetic daily light integral (DLI) and temperature are two environmental factors that profoundly influence plant growth and development. Two common ornamental annual crops, salvia (*Salvia splendens* F. Sello ex Roem & Schult.) and marigold (*Tagetes patula* L.), were grown in glass greenhouses under a mean DLI of 5 to 25 mol·m⁻²·d⁻¹ at temperatures from 14 to 27 °C. Growth (e.g., plant dry weight at flowering) and flowering characteristics (e.g., time to flowering and flower number) were modeled in response to the mean daily temperature and DLI by using multiple regression analysis. The rate of progress to flowering of salvia and marigold was primarily influenced by the mean air temperature. For example, time from seedling transplant to flowering of salvia decreased from 42 days to 24 days as temperature increased from 15 to 25 °C, with a mean DLI of 10 mol·m⁻²·d⁻¹. Flower number and plant dry weight on the date of first flowering generally decreased with increasing temperature and decreasing DLI in both species. For example, marigold plants grown at 15 °C and a mean DLI of 25 mol·m⁻²·d⁻¹ were 2.45 times greater in dry weight, had 2.12 more flowers, and had 49% larger flowers at flowering compared with plants grown at 25 °C and a mean DLI of 5 mol·m⁻²·d⁻¹. The models can be used to predict the impact of changing light and temperature conditions on plant quality and flowering of these two crops.

Commercial production of ornamental annuals is the largest contributor to the value of floriculture in the United States, with a reported wholesale value of $1.9 billion in 2005 (U.S. Department of Agriculture, 2006). The majority of ornamental annual plants are produced in greenhouses from January through May for sales in spring and early summer. During this period, plants are exposed to a wide range of temperature and light conditions. In addition, greenhouse growers often produce multiple crops during this period, so the environmental conditions provided to an early spring crop are usually different from the conditions provided in the same location to a later crop.

Temperature is the most commonly manipulated environmental factor in greenhouse production. The growing temperature used by commercial growers depends on numerous factors, including time of year, greenhouse location, environmental control systems, fuel cost for heating, crop finish dates, desired plant quality and size, stage of development, and plant species. The rate of plant development is a function of temperature and usually increases linearly from above the species-specific base temperature until the optimum temperature (Roberts and Summerfield, 1987). In addition, temperature also influences plant quality parameters such as flower number and size, branching, and mass. For example, flower bud number and flower size of tickseed (*Coreopsis grandiflora* Hogg ex Sweet.), blanket flower (*Gaillardia xgrandiflora* Van Houte), and shasta daisy (*Leucanthemum xsuperbum* Bergman ex. J. Ingram) increased as mean daily temperature decreased from 27 to 15 °C (Yuan et al., 1998).

Ornamental annuals are also exposed to a range of photosynthetic daily light integrals (DLIs), depending on the time of year, greenhouse location, light transmission to the crops, and supplemental lighting. For example, the mean outdoor DLI in Michigan is less than 15 mol·m⁻²·d⁻¹ in January and more than 50 mol·m⁻²·d⁻¹ in May (Korczyński et al., 2002). Assuming a typical greenhouse light transmission of 50% (Hanan, 1998), ornamental annual crops can be grown under a mean DLI less than 8 mol·m⁻²·d⁻¹ to more than 25 mol·m⁻²·d⁻¹. Mean DLI can influence flower initiation and flowering parameters. For example, the number of days required for flower initiation of several cultivars of geranium (*Pelargonium ×domesticum* L.H. Bailey) generally decreased as the DLI increased from 5 to 20 mol·m⁻²·d⁻¹ (Loehrlein and Craig, 2004). The interactive effects of DLI with temperature on plant growth and development have not been described for many floriculture crops.

Ornamental annuals propagated by seed are commonly produced in two distinct stages: the seeding stage and the finish stage. During the seeding stage, seeds are sown in plug trays that have small soil volumes and closely spaced growing cells. Seedlings are raised in plugs until plants have established a sufficient root system and adequate vegetative development, and are deemed ready for transplant. The finish stage begins on the date of transplant into larger containers and continues until plants are marketed. We performed experiments with two annual crops, marigold and salvia, to quantify how temperature and DLI influence growth and development during the finish stage. Marigold and salvia were selected for study because they are among the most popular species produced by the greenhouse industry in the United States (U.S. Department of Agriculture, 2006). Plants were grown at a mean daily temperature of 14 to 27 °C with a mean DLI of 5 to 25 mol·m⁻²·d⁻¹, which represent the range of conditions for most commercial greenhouse
companies that produce ornamental annuals. Plant models were then developed to predict the effect of changing temperature or DLI on growth and development of these two crops.

Materials and Methods

**Seedling plug culture.** Seeds of salvia ‘Vista Red’ and marigold ‘Bonanza Yellow’ were sown in 288-cell (5-mL) plug trays on 25 Jan. and 2 Apr. at a wholesale plug producer (Raker’s Acres, Litchfield, Mich.). The germinated seeds were received at Michigan State University on 29 Jan. and 8 Apr. so that the DLI during the seedling stage could be controlled. The 288-cell trays were placed in a growth chamber set at 23 °C under 150 μmol·m⁻²·s⁻¹ provided by cool-white fluorescent (VHOF96T12; Philips, Bloomfield, N.J.) and incandescent lamps with a 16-h photoperiod. A vapor pressure deficit of 0.7 kPa was maintained. Plugs were top irrigated with acidified well water supplemented with a water-soluble fertilizer to provide micronutrients: 40 mg·L⁻¹ N, 4 mg·L⁻¹ P, 40 mg·L⁻¹ K, and 5 mg·L⁻¹ Ca (Pramuk and Runkle, 2005a). Seedlings were grown until deemed ready for transplant, which was 19 and 26 d from sowing for marigold and salvia respectively.

**Greenhouse temperature and DLI treatments.** For each species and experiment, 150 seedlings were removed from the growth chamber and transplanted into 10-cm-diameter pots (470 mL) containing 70% peatmoss, 21% perlite, and 9% vermiculite (Suremix; Michigan Grower Products, Galesburg, Mich.). Thirty plants of each species were placed (with ≈15-cm center spacing) into five similar glass greenhouse compartments set at constant air temperatures of 14, 17, 20, 23, and 26 °C. In each greenhouse compartment, the water vapor pressure deficit was calculated using a dry and wet bulb temperature and was maintained between 0.7 and 1.0 kPa by steam injection into the air. In the center of each greenhouse section, air temperature was measured by a type E thermocouple (TT-E-40; Omega Engineering, Stamford, Conn.) placed in an aspirated tube. Thermocouples were connected to a CR10 data logger (Campbell Scientific, Logan, Utah), and data were recorded every 10 s.

Three DLI treatments were provided to plants at each of the five temperature treatments, with 10 plants under each DLI and temperature combination. The DLI treatments were delivered using a combination of shade cloth and different light intensities from high-pressure sodium (HPS) lamps from 0600 to 2200 hr to achieve a 16-h photoperiod. A low DLI was provided using ambient light with 50% shade cloth (OLS 50; Ludvig Svensson, Charlotte, N.C.) and supplemental HPS lighting that delivered a photosynthetic photon flux (PPF) of 35 μmol·m⁻²·s⁻¹ at plant level. Plants under the moderate and high DLI treatments were grown under ambient light without shade cloth and with supplemental HPS lighting that provided a PPF of 75 or 170 μmol·m⁻²·s⁻¹ respectively. Line quantum sensors containing 10 photodiodes (Apogee Instruments, Logan, Utah) were placed directly above plants under the three lighting treatments in three greenhouse compartments (nine sensors total) to measure PPF. Sensors were connected to the same CR10 data logger and data were recorded every 10 s.

**Plant culture and data collection.** Plants were top irrigated as necessary with acidified well water [as described by Pramuk and Runkle (2005a)] supplemented with a water-soluble fertilizer containing 125 mg·L⁻¹ N, 13 mg·L⁻¹ P, 125 mg·L⁻¹ K, 15 mg·L⁻¹ Ca, 1 mg·L⁻¹ Fe, 0.1 mg·L⁻¹ B and Mo, and 0.5 mg·L⁻¹ Mn, Zn, and Cu (MSU Special; GreenCare Fertilizers, Kankakee, Ill.). Date of flowering was monitored daily and was recorded, and at flowering, the following were measured: plant height from soil level to the inflorescence apex, number of nodes on the primary shoot, total shoot dry weight, number of inflorescences (flower number), and diameter of the first open floret (salvia) or inflorescence (marigold). Marigold was considered in flower when all petals were fully reflexed; salvia was considered in flower when the first floret opened.

**Data analysis.** Data were analyzed using the calculated mean daily air temperature and DLI for each plant from transplant to the date of flowering. Flowering data were converted to rates (flowering rate) by taking the reciprocal of days to flowering (1/d to flowering). Multiple regression analysis was performed using SAS (SAS Institute, Cary, N.C.) response surface regression (RSREG procedure) to determine the effect of DLI in combination with air temperature. Similar studies with temperature and DLI have used similar forms of analysis (Adams et al., 1997; Carew et al., 2003; Pramuk and Runkle, 2005a). If the contribution of individual terms to the model was nonsignificant at P > 0.05, the terms were removed and the regression (REG procedure) was used to determine the model coefficients. Equations were then used to generate predictive models based on ≈300 observations (both experimental replications) for each species. The models in Fig. 1 and Fig. 2 are in the form of

\[
y = y_0 + aT + bT^2 + cDLI + dDLI^2 + eT \times DLI
\]

where y equals the plant parameter, y₀ is the y-axis intercept, T is temperature (measured in degrees Celsius), DLI is the mean daily light integral (measured in moles per square meter per day), and a, b, c, d, and e are species-specific constants presented in Table 1. Response minimums (e.g., minimum plant height) and maximums (e.g., maximum plant height) were the calculated points of inflection of the response surfaces and were rounded to the nearest 1 °C. Axes for Figs. 1C, 1D, 3C, and 3D were reversed to improve clarity of the data; otherwise, response surfaces would have sloped away, making interpretations more difficult. Base temperature, the temperature at which plant development stops, was estimated under 5 and 15 mol·m⁻²·d⁻¹ by inputting the DLI into the flowering rate equation and setting the equation equal to zero. The DLI parameter was included in the base temperature calculations because plants grown at a high irradiance are usually warmer than plants grown at a lower irradiance at the same air temperature (Pietsch et al., 1995; Shimizu et al., 2004). Therefore, we used air temperature and DLI collectively to estimate the plant temperature at which the flowering rate was zero (base temperature).

**Results**

**Salvia.** The flowering rate was primarily controlled by the mean air temperature, but there was an interaction with mean DLI (Table 1). For example, increasing the temperature from 15 to 25 °C under a mean DLI of 10 mol·m⁻²·d⁻¹ decreased flowering time from 42 d to 24 d; under a mean DLI of 20 mol·m⁻²·d⁻¹, flowering time decreased from 37 d to 21 d (Fig. 1). Flowering rate continued to increase with temperature, and thus an optimum temperature for maximal development was not determined. The flowering time model accurately described the data set of salvia within ± 5 d for 90% of the
actual data (Fig. 2). The calculated base temperature under a mean DLI of 5 or 15 mol m$^{-2}$ d$^{-1}$ was 7.3 or 6.8 °C respectively.

Temperature and DLI did not have a significant effect on node number at flowering, which averaged 8.5 and 7.0 in the January and April experiments respectively (data not presented). Plant height at flowering increased with temperature until a maximum at 20 °C under 5 mol m$^{-2}$ d$^{-1}$ (16.2 cm) and at 24 °C under 25 mol m$^{-2}$ d$^{-1}$ (15.2 cm). Plant height decreased with increasing DLI at temperatures ranging from 14 to 24 °C. Plant height was highly variable, especially at temperatures more than 25 °C, and therefore the $r^2$ value was relatively low.

Flower number generally decreased with increasing temperature, but was between 9 and 11 when the mean daily temperature was ≤ 20 °C. Salvia had the fewest flowers when grown at the warmest temperatures and lowest mean DLI (e.g., plants averaged six flowers when grown at 26 °C and a mean DLI of 6 mol m$^{-2}$ d$^{-1}$). Flower size was not significantly influenced by growing temperature or DLI (data not presented). Shoot dry weight at first flowering was primarily influenced by temperature and increased as temperature decreased, especially when the DLI was low. For example, mean dry weight of salvia was 42% greater when grown at 18 °C than at 24 °C at a mean DLI of 10 mol m$^{-2}$ d$^{-1}$.

**MARIGOLD.** Flowering rate increased as DLI and temperature increased, and a temperature for maximal development was not determined (Table 1; Fig. 3). The flowering time model accurately described the data set within ± 5 d for 91% of the actual data (Fig. 2). Plants grown under a mean DLI of 10 mol m$^{-2}$ d$^{-1}$ flowered in 36 d at 15 °C and in 23 d at 25 °C. Increasing the DLI to 20 mol m$^{-2}$ d$^{-1}$ reduced flowering time by 6 d at 15 °C and 4 d at 25 °C. The calculated base temperature was –2.9 °C, regardless of DLI. The number of nodes that developed before flowering was not influenced by temperature or mean DLI and averaged 5.4 (data not shown).

Mean plant height at flowering increased by 0.17 cm for every 1 °C increase in average daily temperature and by 0.07 cm for every 1 mol m$^{-2}$ d$^{-1}$ increase in the DLI. Flower number and flower diameter were greatest when DLI was high and the average temperature was low. Within the DLI range tested, an increase in DLI by 5 mol m$^{-2}$ d$^{-1}$ increased flower number by 1.3 and flower diameter by 0.16 cm. Lowering the mean daily temperature by 5 °C increased flower number at first flowering by 3.7 and increased flower size by 0.65 cm. Dry weight was greatest at the coolest temperatures and highest DLI, and decreased with increasing temperature. Plants grown at 15 °C and a DLI of 25 mol m$^{-2}$ d$^{-1}$ were 2.45 times greater in

![Fig. 1. (A–D) Temperature and daily light integral (DLI) effects on salvia flowering rate (A: 1/d to flowering) and plant height (B), flower bud number (C), and shoot dry weight at flowering (D). The coefficients for the models are presented in Table 1. Symbols represent measured data, and response surfaces represent model predictions. Axes for C and D were reversed to improve clarity of data.](image)
shoot dry weight, had 2.12 more flowers, and had 49% larger flowers than plants grown at 25 °C and a mean DLI of 5 mol·m⁻²·d⁻¹.

**Discussion**

The flowering rate of salvia and marigold was primarily controlled by temperature within the experimental conditions provided. Similarly, mean DLI during the finish stage did not influence node number at flowering in celosia (*Celosia argentea* L. var. *plumosa* L.) and impatiens (*Impatiens wallerana* Hook.f.) (Pramuk and Runkle, 2005a). Faust et al. (2005) reported that increasing the DLI from 5 to 43 mol·m⁻²·d⁻¹ during the finish stage accelerated flowering (by <5 d) in *Salvia coccinea* L., but *Tagetes erecta* L. flowered at a similar time under a DLI of 19 or 43 mol·m⁻²·d⁻¹; node number at flowering was not reported. In contrast, an increase in DLI during the seedling stage accelerated subsequent flower initiation in the same salvia and marigold cultivars used in this study as well as in celosia and impatiens (Pramuk and Runkle, 2005b). This suggests that plants used in this study had at least partially initiated flowers during the seedling stage and before the onset of treatments, or that the promotive effects of a high DLI on flower initiation decrease with an increase in plant size or maturity.

Flowering rate of marigold and salvia continued to increase with mean daily temperature up to the warmest treatment (27 °C), and thus a maximum developmental rate was not identified for either species. In marigold, whole-plant net photosynthesis was greater in plants grown at 30 °C than at 20 °C, which suggests the maximal developmental rate is achieved at more than 30 °C (van Iersel and Seymour, 2003). Temperatures for maximal rates of development have been reported for several other bedding plant species, including pansy (*Viola × wittrockiana* Gams.; 22 °C), dahlia (*Dahlia pinnata* Cav.; 24 °C), petunia (*Petunia × hybrida* Vilm.-Andr.; 25 °C), celosia (25 °C), impatiens (26 °C), and vinca (*Catharanthus roseus* L., 35 °C) (Adams et al., 1997; Brøndum and Heins, 1993; Kaczperski et al., 1991; Pietsch et al., 1995; Pramuk and Runkle, 2005a).

**Table 1. Parameters of regression analysis relating flowering rate, plant height, shoot dry weight, and node and flower number for salvia and marigold to mean air temperature (°C) and daily light integral (DLI measured in mol·m⁻²·d⁻¹).**

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Salvia</th>
<th>Marigold</th>
<th>r²</th>
<th>SE</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Flowering rate (1/d to flowering)</td>
<td>-2.41 E–2</td>
<td>1.15 E–3</td>
<td>0.84</td>
<td>1.40</td>
<td>0.05***</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>4.45 (2.84)</td>
<td>1.61 (0.49)</td>
<td>0.83</td>
<td>0.50</td>
<td>0.02***</td>
</tr>
<tr>
<td>Dry weight (g)</td>
<td>14.02 (0.69)</td>
<td>1.28 (0.36)</td>
<td>0.94</td>
<td>0.20</td>
<td>0.00***</td>
</tr>
<tr>
<td>Flowers (n)</td>
<td>14.02 (0.69)</td>
<td>1.28 (0.36)</td>
<td>0.94</td>
<td>0.20</td>
<td>0.00***</td>
</tr>
<tr>
<td>Flower diameter (cm)</td>
<td>7.00 (0.13)</td>
<td>3.17 (0.05)</td>
<td>0.97</td>
<td>0.03</td>
<td>0.00***</td>
</tr>
<tr>
<td>Coefficients for model equations were used to generate Figs. 1 and 2.</td>
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All models are in the form of $y = y + aT + bT^2 + cDLI + dDLI^2 + eT·DLI$. 

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Fig. 2. Frequency of predicted minus actual days to flowering in salvia and marigold grown at different temperatures (≈14 to 27 °C) and daily light integrals (≈5 to 25 mol·m⁻²·d⁻¹). The models used to predict flowering time are shown in Table 1 and are based on more than 290 observations for each species.
The base temperature of a plant is usually identified by extrapolating the rate of development at temperatures below the data set until the rate reaches zero, and assumes linearity. The base temperatures calculated for marigold and salvia were based on the effect of air temperature and DLI on plant temperature, although DLI had little or no effect on the estimates for marigold (\(-3\) °C) and salvia (\(\approx 7\) °C). In contrast, the estimated base temperature of celosia and marigold decreased by 1.5 °C and 3.2 °C respectively, as DLI increased from 5 to 15 mol m\(^{-2}\) d\(^{-1}\) (Pramuk and Runkle, 2005a). The base temperature has been calculated for other herbaceous ornamental plants without consideration of DLI, including shasta daisy (\(-3.4\) °C), gaillardia (3.3 °C), dahlia (5.5 °C), and tickseed (6.8 °C) (Brøndum and Heins, 1993; Yuan et al., 1998).

Although the flowering rate increased with temperature in both species, plant quality parameters decreased, especially when the DLI was low. As daily temperature increased from 15 to 25 °C, flower number decreased from 10.3 to 7.0 in salvia and from 19.7 to 12.4 in marigold when the mean DLI was 10 mol m\(^{-2}\) d\(^{-1}\). In addition, flower size of marigold decreased by 24%. Under the same DLI, models predicted the same 10 °C increase in forcing temperature to decrease flower number of celosia by 18% and impatiens by 58% (Pramuk and Runkle, 2005a). In pansy, flower size decreased by 26% or 37% as temperature increased from 18 to 24 °C under a mean DLI of 15.6 mol m\(^{-2}\) d\(^{-1}\) or 4.1 mol m\(^{-2}\) d\(^{-1}\) respectively (Niu et al., 2000). Similarly, flower diameter of dahlia decreased by 1.4 mm°C\(^{-1}\) as temperature increased from 11 to 30 °C (Brøndum and Heins, 1993). The effects of forcing temperature on flower number and size have also been described in several herbaceous perennials. For example, an increase in forcing temperature from 16 to 26 °C decreased flower number of tickseed, rudbeckia (\textit{Rudbeckia fulgida} Ait. ‘Goldsturm’), and shasta daisy by 80%, 75%, and 55% respectively (Yuan et al., 1998). In addition, a decrease in forcing temperature from 26 to 14 °C increased flower size of campanula (\textit{Campanula carpatica} Jacq.) and \textit{Campanula} L. ‘Birch Hybrid’ by 35% (Niu et al., 2001).

The increase in flower number of salvia (when temperature was \(\geq 19\) °C) and flower number and size of marigold from an increasing DLI is consistent with data for several other greenhouse crops, including wax begonia (\textit{Begonia ×semperiflora-cultorum} L.), campanula, celosia, hibiscus (\textit{Hibiscus radiatus} Cav.), impatien, marigold, pansy, petunia, vinca (\textit{Catharanthus roseus} L.), and zinnia (\textit{Zinnia elegans} L.) (Faust et al., 2005; Niu et al., 2000, 2001; Pramuk and Runkle, 2005a).
2005a; Warner and Erwin, 2003). However, few studies have quantified the interactive effects of temperature and DLI on flowering characteristics. In *Primula vulgaris* Huds., the DLI that elicited the most rapid flower initiation decreased (to 11 mol·m⁻²·d⁻¹) as temperature increased to 13 °C, then increased with temperature (Karlsson, 2002). Our data indicate that at a warmer growing temperature, a higher DLI is required to produce a plant of comparable quality to plants grown at a cooler temperature. For example, to produce a crop of marigold with ≥15 flower buds, the model indicates that plants could be grown at a mean daily temperature of less than 20 °C when the DLI is 5 mol·m⁻²·d⁻¹, but at 25 °C, a mean DLI ≥ 20 mol·m⁻²·d⁻¹ is required.

Time from transplant to flower increased with decreasing temperature, and thus plants grown at the lower temperatures were older and had a greater accumulated DLI before flowering. Dry weight of marigold increased as DLI increased from 5 to 25 mol·m⁻²·d⁻¹, but the effect of DLI on salvia depended on temperature. Salvia grown at 26 °C had the greatest dry weight under a DLI of 22 mol·m⁻²·d⁻¹, whereas dry weight was greatest at 16 °C when the DLI was 13 mol·m⁻²·d⁻¹. In celosia and impatiens, dry weight at flowering increased as DLI increased from 5 to 25 mol·m⁻²·d⁻¹, regardless of temperature (Pramuk and Runkle, 2005a). Faust et al. (2005) reported increases in dry weight with increasing DLI for several bedding plants grown outdoors at a mean daily temperature of ≈23 °C. However, shoot dry weight of the shade-tolerant wax begonia and impatiens did not increase when the DLI increased from 19 to 43 mol·m⁻²·d⁻¹. Nameli and van Iersel (2004) reported that dry weight of wax begonia increased, but at a decreasing rate as DLI increased from 5.3 to 19.4 mol·m⁻²·d⁻¹.

The models generated can be used to predict the effects of temperature and DLI on growth and flowering of marigold and salvia in commercial settings. For example, a salvia crop grown at 20 °C in Michigan and Florida in March could receive a mean DLI of 12 and 20 mol·m⁻²·d⁻¹ respectively, assuming 50% greenhouse light transmission (Korzynski et al., 2002). The models predict that salvia would take about 3 d longer to flower in Michigan, but dry weight and flower number would be essentially the same in both locations. In the same scenario, marigolds grown in Florida would flower 4 d earlier, have 5% larger flowers, 12% more flower buds, and 23% greater dry weight compared with a crop grown under the lower DLI in Michigan that time of year. In addition, the data illustrate that plants of similar quality can be grown at a low mean daily temperature and DLI as at a high temperature and high DLI. For example, the flower number model for marigold predicts 17.7 buds per plant if grown at 17 °C and a DLI of 8 mol·m⁻²·d⁻¹ or at 23 °C and a DLI of 25 mol·m⁻²·d⁻¹. However, marigold grown at 17 °C would take 15 d longer to flower than at 23 °C. The models were developed with long-day conditions, and for photoperiodic salvia cultivars, the model may be less predictable if plants are grown under short days. Additional studies are merited to determine the effects of temperature and DLI on growth and development of other economically important greenhouse crops.

**Literature Cited**


