



Article Supplemental Lighting Quality Influences Time to Flower and Finished Quality of Three Long-Day Specialty Cut Flowers

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Abstract: Year-round demand for locally sourced specialty cut flowers continues to increase. However, due to low radiation intensities and temperatures, growers in northern latitudes must utilize greenhouses, but limited production information detailing manipulation of the radiation environment exists. Therefore, our objective is to quantify the influence of supplemental lighting (SL) quality on time to flower and harvest and stem quality of three long-day specialty cut flowers. Godetia 'Grace Rose Pink' (Clarkia amoena), snapdragon 'Potomac Royal' (Antirrhinum majus), and stock 'Iron Rose' (Matthiola incana) plugs are transplanted into bulb crates and placed in one of six greenhouse compartments with SL providing a total photon flux density of 120 μ mol·m⁻²·s⁻¹ from 0700 to 1900 HR. After four weeks, SL is extended to provide a 16 h photoperiod to induce flowering. SL treatments are provided by either high-pressure sodium (HPS) fixtures or various light-emitting diode (LED) fixtures. Treatments are defined by their 100 nm wavebands of blue (B; 400-500 nm), green (G; 500-600 nm), red (R; 600-700 nm), and far-red (FR; 700-800 nm) radiation (photon flux density in μ mol·m⁻²·s⁻¹) as B₇G₆₀R₄₄FR₉ (HPS₁₂₀), B₂₀G₅₀R₄₅FR₅, B₂₀R₈₅FR₁₅, B₃₀G₂₅R₆₅, B₁₂₀, or R₁₂₀. Time to harvest (TTH) is up to 14, 15, and 10 d slower under R_{120} SL for godetia, snapdragon, and stock, respectively, compared to the quickest treatments (HPS₁₂₀, B₁₂₀, and B₂₀R₈₅FR₁₅ SL). However, R₁₂₀ SL produces cut flowers up to 18% longer than those grown under the quickest treatments. Both broad-spectrum LED fixtures slightly delay TTH compared to the quickest treatments. Stem caliper is not commercially different between treatments for godetia or snapdragon, although stems are up to 14% thinner for stock grown under B120 SL compared to the other treatments. Flower petal color is not commercially different between SL treatments. We recommend utilizing a SL fixture providing a spectrum similar to $B_{20}R_{85}FR_{15}$ SL or $B_{20}G_{50}R_{45}FR_5$, as they elicit desirable crop responses with minimal developmental, quality, and visibility tradeoffs. While HPS lamps perform similarly to the recommended fixtures, we recommend utilizing LEDs for their higher photon efficacy and potential energy savings.

Keywords: high-pressure sodium lamps; light-emitting diodes; light quality; controlled-environment agriculture; greenhouse

1. Introduction

Demand for locally produced specialty cut flowers exists year-round [1,2]. However, low radiation intensities and temperatures in northern latitudes prohibit the production of specialty cut flowers outdoors or in unheated high tunnels during the winter and early spring [3,4]. For example, the outdoor solar daily light integral (DLI) can fall to as low as 5 to $10 \text{ mol} \cdot \text{m}^{-2} \cdot \text{d}^{-1}$ [5] and as low as $\leq 5 \text{ mol} \cdot \text{m}^{-2} \cdot \text{d}^{-1}$ in controlled-environment greenhouses because of reflection from glazing and shading from the superstructure [6]. Because of these unfavorable environmental conditions, greenhouses with high-intensity supplemental lighting (SL) must be employed to maintain environmental conditions suitable for cut flower growth, so growers can tap into local markets and satisfy consumer demand throughout the year.



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In recent years, the advent of horticultural light-emitting diodes (LEDs) has given growers the potential to further customize the emission spectra of their lighting sources, allowing for the inclusion of narrow wavebands [7–10]. Thus, a large variety of SL fixtures with different static or customizable emission spectra have become commercially available. The composition of radiation emitted from a lighting fixture can have substantial effects on plant growth and development, especially when the solar DLI is low [11,12], with some wavebands acting not only as photosynthetic stimuli, but as developmental signals [13]. Photosynthetically active radiation (PAR; 400–700 nm) is primarily responsible for driving photosynthesis, although isolated wavebands within and outside of this range can bring about specific photomorphogenic responses. Although outside of the traditional definition of PAR, far-red (FR) radiation (700-800 nm) has recently been shown to contribute to photosynthesis directly by working synergistically with photons within the traditional designation of PAR, and indirectly by promoting leaf expansion [14–18]. However, the inclusion of FR radiation in the range of PAR is yet to be widely accepted by the greater scientific community. For decades, however, it has been broadly understood that FR radiation predominantly influences plant morphology and development [9,17,19,20].

Photomorphogenic responses such as internode elongation, leaf expansion, and flowering are regulated by various photoreceptors within plant cells including cryptochromes, phototropins, and phytochromes [10,12,17]. For instance, a decreasing ratio of red (R; 600–700 nm) and FR radiation emitted from a radiation source generally promotes extension growth [9,21], which is a function of phytochrome photoreceptors [21,22]. The influence of R and FR radiation on crop morphology is well-documented. For instance, Elkins and van Iersel [19] reported that the height of foxglove 'Dalmatian Peach' (*Digitalis purpurea*) cut flower seedlings grown under sole-source lighting for 16 h·d⁻¹ increased by 38% as the R to FR ratio of the light source decreased from 13.7 to 0.6.

Phytochrome photoreceptors exist in two reversible conformations: P_R and P_{FR} . These conformations are designated as the "inactive" and "active" conformations, respectively [10,23], as P_{FR} is primarily responsible for initiating phytochrome-mediated photomorphogenic responses on the cellular level [12]. The ratio of R:FR radiation in a radiation source's spectrum can influence the ratios of these conformations. When exposed to R radiation, P_R changes conformation to P_{FR} , while P_{FR} can revert to P_R in the presence of either FR radiation or through natural degradation [24]. The ratio of these phytochrome conformations is referred to as the phytochrome photoequilibrium (PPE; P_{FR}/P_{R+FR}), and it is closely associated with the activity of phytochromes within plant cells [23,24].

R and FR radiation are not only prominent drivers of crop architecture; they are also integral to the flowering responses of many long-day plants (LDPs) [25,26]. When grown under a FR-radiation deficient filter, flowering of campanula 'Blue Clips' (*Campanula carpatica*), coreopsis 'Early Sunrise' (*Coreopsis* × grandiflora) and pansy 'Crystal Bowl Yellow' (*Viola* × wittrockiana) was delayed by 2, 14, and 21 d, respectively, compared to plants grown under a neutral filter that allowed for the transmission of FR radiation [26]. It has also been shown that SL emitting moderate intensities of FR radiation (\geq 15 µmol·m⁻²·s⁻¹) can hasten flowering in LDPs compared to SL without FR radiation [27]. For instance, the LDP snapdragon 'Liberty Classic Yellow' (*Antirrhinum majus*) grown under SL containing 15 µmol·m⁻²·s⁻¹ of FR radiation for a 16 h·d⁻¹ for 28 d during the plug stage reached open flower 6 d faster than plants grown under SL containing only blue (B) and R radiation during the plug stage [27].

B radiation (400–500 nm) inhibits extension growth in many crops, which is a function of cryptochrome and phototropin photoreceptors [28–30]. However, B radiation mediated stem compaction responses are species-specific, and some crops defy this phenomenon [12]. In a 2017 study, Poel and Runkle reported that geranium 'Pinto Premium Salmon' (*Pelargonium ×hortorum*) and petunia 'Single Dreams White' (*Petunia ×hybrida*) grown under (%) B₄₅R₅₅ LEDs emitting a photosynthetic photon flux density (PPFD) of $90 \pm 10 \ \mu\text{mol} \cdot \text{m}^{-2} \cdot \text{s}^{-1}$ for 16 h·d⁻¹ were ≈17% and 22% shorter, respectively, than those grown under SL provided by B₁₀G₅R₈₅ LEDs [10]. In a separate study, poinsettia 'Christmas Spirit' and 'Christmas Eve' (*Euphorbia pulcherrima*) grown under 100 ± 20 μ mol·m⁻²·s⁻¹ of high-pressure sodium (HPS) SL with 5% B radiation for 10 $h \cdot d^{-1}$ were \approx 52% and 36% taller, respectively, than those grown under the same intensity and duration of SL provided by LEDs emitting 20% B radiation for 12 weeks [31].

Additionally, a moderate intensity of B radiation can function as a long-day signal for some crops. For instance, a 4 h night interruption (NI) provided by 30 μ mol·m⁻²·s⁻¹ of B radiation was as effective as a 4 h NI provided by 2 μ mol·m⁻²·s⁻¹ from R + white (W) + FR LEDs at promoting flowering in calibrachoa 'Callie Yellow Improved' (*Calibrachoa × hybrida*), coreopsis 'Early Sunrise', petunia 'Wave Purple Improved', rudbeckia 'Indian Summer' (*Rudbeckia hirta*), and snapdragon 'Liberty Classic Yellow' [32]. Furthermore, Sharath Kumar et al. [33] demonstrated the efficacy of a 4 h day extension (DE) provided by 40 μ mol·m⁻²·s⁻¹ of 100% B radiation at inhibiting flowering of greenhouse-grown chrysanthemum 'Radost' (*Chrysanthemum morifolium*).

Traditionally, high-intensity horticultural LED fixtures utilized a combination of B and R diodes because of the higher absorption of B and R photons in upper leaf cells, consistent with the peak absorbances of chlorophyll a and b, compared to other wavebands [34,35]. However, recent research has found that green radiation (G; 500–600 nm) can be comparably effective for photosynthesis. For example, Liu and van Iersel [35] reported that whole-plant photosynthetic efficacy of G radiation applied to lettuce 'Green Towers' (*Lactuca sativa*) was higher than that of B radiation when applied at intensities > 500 μ mol·m⁻²·s⁻¹, as G photons are transmitted farther into the plant canopy than other wavebands [8,35].

In addition to stimulating photosynthesis, G radiation has been shown to inhibit branching of some ornamental plants when applied at moderate intensities [36]. For example, petunia 'Easy Wave Burgundy Star' had an average of roughly five fewer lateral branches when the G radiation photon flux density (PFD) during a 16 h DE was 25 μ mol·m⁻²·s⁻¹ compared to 2 μ mol·m⁻²·s⁻¹. Furthermore, moderate fluxes of G radiation can serve as a long-day signal for some floriculture crops [36]. G radiation saturated the flowering response of ageratum 'Hawaii Blue' (Ageratum houstonianum) when applied at intensities of 2 μ mol·m⁻²·s⁻¹ during a 16 h DE, although 13 μ mol·m⁻²·s⁻¹ was required to saturate the flowering responses of petunia 'Easy Wave Burgundy Star' and 'Wave Purple Improved', and snapdragon 'Liberty Classic Yellow' [36]. Furthermore, in several studies on nonhorticultural crops, G radiation inhibited some B radiation-mediated photomorphogenic responses, such as hypocotyl compaction and anthocyanin accumulation [12]. In addition, certain fluxes of G radiation can elicit stem elongation responses similar to that of FR radiation, which can be counteracted with B radiation [37]. Interestingly, stem elongation of plants exposed to a combination of G and FR radiation was greater than that of plants exposed to either waveband alone [38].

When applied simultaneously, B, G, R, and FR wavebands can have compounding effects on crop growth and development. Height of high-wire cucumber 'Elsie' (*Cucumis sativus*) and tomato 'Climstar' (*Solanum lycopersicum*) were up to $\approx 17\%$ and 25% taller when grown under 120 µmol·m⁻²·s⁻¹ of B₃₀G₃₀R₆₀ SL for 16 h·d⁻¹ compared to the same intensity and duration of B₂₅R₉₅ SL, suggesting that the addition of G radiation counteracted B-mediated plant compaction, producing taller plants [39]. Moreover, when B, G, and R radiation is applied together, the resulting broad-spectrum radiation appears white (W) to the human eye, increasing the visibility in the work environment. This can aid in detection of pests and nutrient deficiencies compared to spectra comprised of one or two wavebands [34].

Radiation quality can also influence flower petal color by influencing the accumulation of pigments such as anthocyanins, carotenoids, and flavonoids [40,41]. Petal color is influenced in part by petal morphology, i.e., tissue thickness and inhomogeneity [40], which may be affected by radiation quality. While flower color is of ecological importance to angiosperms as it helps attract specific pollinators [40], it is also of significant aesthetic importance to consumers [42]. Manipulating radiation quality to produce cut flowers with more vibrant colors can increase consumers' willingness to buy and the subsequent product enjoyment [42].

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To our knowledge, minimal research examining the influence of supplemental radiation quality on the greenhouse production of LDP specialty cut flowers has been published and, thus, additional research could provide utility to cut flower greenhouse growers. Therefore, the objective of this study was to quantify the influence of SL radiation quality on time to flower and harvest and on the finished quality of three long-day specialty cut flowers. We hypothesized that flowering would be delayed for plants grown under SL lacking FR radiation. Additionally, we hypothesized that plants grown under R₁₂₀ SL would exhibit greater stem elongation compared to the other treatments, particularly B₁₂₀ SL, where we predicted that stems would remain compact. We also postulated that treatments with a combination of B, G, R, and FR wavebands would yield shorter cut flowers when the emission spectrum contained a higher flux of B radiation and longer cut flowers when the emission spectrum contained a higher flux of FR radiation.

2. Materials and Methods

2.1. Plant Material, Culture, and Lighting Treatments

Seeds of godetia 'Grace Rose Pink' (*Clarkia amoena*; Sakata Seed America, Morgan Hill, CA, USA), snapdragon 'Potomac Royal' (PanAmerican Seed, West Chicago, IL, USA), and stock 'Iron Rose' (*Matthiola incana*; Sakata Seed America) were sown in 162-cell trays at a commercial propagator (Raker-Roberta's Young Plants, Litchfield, MI, USA). Three trays each of godetia, snapdragon, and stock were received one day after sowing on 18 December 2020 (Replication (Rep.) 1) and 28 December 2021 (Rep. 2).

Young plants were grown in a glass-glazed greenhouse under a natural short-day photoperiod with LED fixtures (Philips GP-TOPlight DRW-MB; Koninklijke Philips N.V., Eindhoven, The Netherlands) providing a supplemental PPFD of $\approx 200 \ \mu mol \cdot m^{-2} \cdot s^{-1}$ from 0730 to 1730 HR, creating a DLI of $\approx 15 \ mol \cdot m^{-2} \cdot d^{-1}$. The greenhouse air average daily temperature (ADT) set point during young plant culture was a constant 16 °C. Stock were thinned after cotyledon expansion to increase the amount of double flowering phenotypes, according to protocols provided by the breeder [43]. Godetia and snapdragon were thinned upon cotyledon expansion. Young plants were irrigated as needed with MSU Plug Special (13 N–2.2 P–10.8 K water-soluble fertilizer containing (mg·L⁻¹) 61 nitrogen, 10 phosphorus, 50 potassium, 28.1 calcium, 4.7 magnesium, 1.3 iron, 0.6 manganese, 0.6 zinc, 0.6 copper, 0.4 boron, and 0.1 molybdenum (GreenCare Fertilizers Inc., Kankakee, IL, USA)) blended with reverse-osmosis water and applied with a mist nozzle (Super Fine Fogg-It Nozzle; Fogg-It Nozzle; Fogg-It Nozzle Co. Inc., Belmont, CA, USA).

After 30 d under short days (17 January 2021 (Rep. 1) and 27 January 2022 (Rep. 2)), 180 godetia, snapdragon, and stock young plants were randomly selected for transplant. Seventy-two bulb crates (39.3 cm wide \times 59.7 cm long \times 17.8 cm tall; 0.04 m³) were filled with a soilless medium containing (by volume) 70% peat moss, 21% perlite, and 9% vermiculite (Suremix; Michigan Grower Products Inc., Galesburg, MI, USA). Each bulb crate held 10 young plants of an individual genus, yielding 18 total bulb crates per genus. Young plants were transplanted at a density of 43 plants per m².

Three bulb crates of each genus were placed on benches on the ground in one of six glass-glazed greenhouse compartments. High-intensity SL fixtures providing a total photon flux density of 120 μ mol·m⁻²·s⁻¹ from 0700 to 1900 HR, creating a total DLI of $\approx 11 \text{ mol·m}^{-2} \cdot d^{-1}$. This was denoted as the vegetative stage. After four weeks, SL duration was increased to provide a 16 h photoperiod from 0600 to 2200 HR, creating a total DLI of $\approx 15 \text{ mol·m}^{-2} \cdot d^{-1}$. This was denoted as the reproductive stage. Whitewash (KoolRay Classic Liquid Shade, Continental Products, Euclid, OH, USA) and/or opaque black cloth covered compartment walls to prevent radiation pollution between compartments and adjacent greenhouses. A quantum sensor (LI-190R, LI-COR Biosciences, Lincoln, NE, USA) positioned horizontally at plant height in each compartment measured the PPFD every 10 s and a datalogger (CR1000; Campbell Scientific, Logan, UT, USA) recorded hourly averages. The actual DLIs during the vegetative and reproductive stages of the two replications of the experiment were calculated and are provided in Tables 1 and 2.

Table 1. Actual daily light integrals (DLIs) (mean \pm SD (mol·m⁻²·d⁻¹)), average daily temperatures (ADTs), mean day temperature, mean night temperature, and mean leaf temperature (mean \pm SD (°C)) for each supplemental light (SL) treatment during the vegetative (VEG) and reproductive (REP) stages of replication 1. SL treatments consisted of either 460 W HPS fixtures (HPS₁₂₀; LR48877; P.L. Light Systems, Beamsville, ON, Canada), 631 W LED fixtures (B₂₀G₅₀R₄₅FR₅; VYPR 2p; Fluence, Austin, TX, USA), 325 W LED fixtures (B₂₀R₈₅FR₁₅; LumiGrow Pro 325; LumiGrow, Emeryville, CA, USA), 600 W LED fixtures (B₃₀G₂₅R₆₅; LX601G, Heliospectra, Göteborg, Sweden), a combination of 72 W LED fixtures (B₁₂₀; HortiLED MULTI, P.L. Light Systems) and 625 W LED fixtures (R₁₂₀; LumiGrow Pro 650E, LumiGrow), or 625 W LED fixtures (LumiGrow Pro 650E; LumiGrow).

SL Treatment and Stage	$\begin{array}{l} \textbf{DLI}\\ \textbf{(Mean}\pm\textbf{SD}\\ \textbf{(mol}{\cdot}\textbf{m}^{-2}{\cdot}\textbf{d}^{-1}\textbf{)} \end{array}$	ADT (Mean \pm SD (°C))	Day Temperature (Mean \pm SD (°C))	Night Temperature (Mean \pm SD (°C))	Leaf Temperature (Mean \pm SD (°C))
HPS ₁₂₀					
VEG	10.7 ± 2.0	15.6 ± 0.6	18.4 ± 1.6	12.7 ± 0.8	17.9 ± 3.5
REP	15.7 ± 4.5	16.1 ± 1.9	18.6 ± 2.7	13.5 ± 3.2	18.3 ± 3.4
B20G50R45FR5					
VEG	10.8 ± 2.0	15.6 ± 0.5	18.3 ± 0.8	12.9 ± 1.0	17.0 ± 2.7
REP	15.6 ± 5.0	16.2 ± 1.7	18.9 ± 2.5	13.4 ± 3.1	18.4 ± 3.6
$B_{20}R_{85}FR_{15}$					
VEG	10.6 ± 2.0	15.6 ± 0.4	18.2 ± 0.8	12.9 ± 0.7	17.1 ± 2.9
REP	15.4 ± 8.9	16.2 ± 1.8	18.8 ± 2.5	16.2 ± 1.8	19.9 ± 4.4
$B_{30}G_{25}R_{65}$					
VEG	10.9 ± 2.1	15.5 ± 0.4	18.1 ± 0.8	12.9 ± 0.6	17.6 ± 2.9
REP	15.1 ± 4.2	16.2 ± 1.8	18.9 ± 2.6	13.5 ± 3.2	18.3 ± 2.7
B ₁₂₀					
VEG	10.8 ± 2.1	16.1 ± 0.5	18.8 ± 1.3	13.3 ± 0.8	17.6 ± 0.9
REP	15.0 ± 4.1	16.5 ± 1.6	19.3 ± 2.5	13.7 ± 2.9	18.8 ± 2.8
R ₁₂₀					
VEG	11.5 ± 2.5	15.5 ± 0.8	18.4 ± 2.6	12.6 ± 0.9	16.5 ± 3.8
REP	15.9 ± 5.3	16.0 ± 1.8	18.3 ± 3.1	13.8 ± 2.8	17.9 ± 3.5

SL treatments consisted of either 460 W HPS fixtures (LR48877; P.L. Light Systems, Beamsville, ON, Canada), 631 W LED fixtures (VYPR 2p; Fluence, Austin, TX, USA), 325 W LED fixtures (LumiGrow Pro 325; LumiGrow, Emeryville, CA, USA), 600 W LED fixtures (LX601G, Heliospectra, Göteborg, Sweden), a combination of 72 W LED fixtures (HortiLED MULTI, P.L. Light Systems) and 625 W LED fixtures (LumiGrow Pro 650E, LumiGrow), or 625 W LED fixtures (LumiGrow Pro 650E; LumiGrow). SL treatments, defined by the PFD delivered at each 100 nm waveband of B (400–500 nm), G (500–600 nm), R (600–700 nm), and FR (700–800 nm) radiation, were $B_7G_{60}R_{44}FR_9$ (HPS₁₂₀), $B_{20}G_{50}R_{45}FR_5$, $B_{20}R_{85}FR_{15}$, $B_{30}G_{25}R_{65}$, B_{120} , or R_{120} , respectively. The spectral distribution of the SL fixtures was measured at crop height in ten random locations throughout each compartment with a spectrometer (LI-180; LI-COR Biosciences) and are presented in Figure 1. The PPE of each SL treatment were estimated according to Sager et al. (1988) and are presented in Table 3.

Two layers of 15 cm supportive netting (HGN32804; Hydrofarm, Petaluma, CA, USA) were positioned \approx 15 and 30 cm, respectively, above each bench. Greenhouse compartments were equipped with evaporative-pad cooling and radiant hot water heating, which, in addition to lighting fixtures, were controlled by an environmental control system (Priva Office version 725–3030, Vineland Station, ON, Canada). The air ADT set points in each greenhouse compartment were 15.8 °C (day/night 18.5/13 °C), with day temperatures maintained from 0800 to 1900 HR and night temperatures maintained from 1900 to 0800 HR. An aspirated thermocouple (36-gauge (0.127 mm diameter) type E, Omega Engineering, Stamford, CT, USA) positioned in the middle of each compartment measured the air temperature at plant height every 10 s, and the datalogger recorded hourly means. Additionally, an infrared thermocouple (Type T, OS36-01; Omega Engineering) positioned against an individual leaf of a snapdragon plant in each compartment measured leaf temperature every 10 s, and the datalogger recorded hourly means. The actual air ADTs, average

daytime and nighttime temperatures at plant height, as well as average leaf temperatures of each treatment during the vegetative and reproductive stages of the two reps. of the experiment were calculated and are provided in Tables 1 and 2.

Table 2. Actual daily light integrals (DLIs) (mean \pm SD (mol·m⁻²·d⁻¹)), average daily temperatures (ADTs), mean day temperature, mean night temperature, and mean leaf temperature (mean \pm SD (°C)) for each supplemental light (SL) treatment during the vegetative (VEG) and reproductive (REP) stages of replication 2. SL treatments consisted of either 460 W high-pressure sodium fixtures (HPS₁₂₀; LR48877; P.L. Light Systems, Beamsville, ON, Canada), 631 W light-emitting diode (LED) fixtures (B₂₀G₅₀R₄₅FR₅; VYPR 2p; Fluence, Austin, TX, USA), 325 W LED fixtures (B₂₀R₈₅FR₁₅; LumiGrow Pro 325; LumiGrow, Emeryville, CA, USA), 600 W LED fixtures (B₃₀G₂₅R₆₅; LX601G, Heliospectra, Göteborg, Sweden), a combination of 72 W LED fixtures (B₁₂₀; HortiLED MULTI, P.L. Light Systems) and 625 W LED fixtures (R₁₂₀; LumiGrow Pro 650E, LumiGrow), or 625 W LED fixtures (LumiGrow Pro 650E; LumiGrow).

SL Treatment and Stage	$\begin{array}{c} \text{DLI}\\ (\text{Mean}\pm\text{SD}\\ (\text{mol}\cdot\text{m}^{-2}\cdot\text{d}^{-1}))\end{array}$	ADT (Mean \pm SD (°C))	Day Temperature (Mean \pm SD (°C))	Night Temperature (Mean \pm SD (°C))	Leaf Temperature (Mean \pm SD (°C))
HPS ₁₂₀					
VEG	11.7 ± 2.2	16.0 ± 0.5	18.7 ± 2.1	13.3 ± 0.8	18.0 ± 4.0
REP	16.1 ± 4.6	16.0 ± 1.0	18.9 ± 2.1	12.9 ± 1.5	18.7 ± 4.0
$B_{20}G_{50}R_{45}FR_5$					
VEG	10.9 ± 2.8	16.3 ± 0.7	19.0 ± 2.4	13.6 ± 0.8	18.0 ± 3.1
REP	16.2 ± 5.2	15.9 ± 0.8	18.9 ± 2.2	12.9 ± 1.1	18.3 ± 3.2
B20R85FR15					
VEG	11.6 ± 2.6	16.0 ± 0.5	18.7 ± 2.1	13.3 ± 0.8	18.3 ± 3.9
REP	15.9 ± 3.9	15.9 ± 1.0	18.9 ± 2.4	12.9 ± 1.3	18.5 ± 3.7
$B_{30}G_{25}R_{65}$					
VEG	11.6 ± 2.6	16.3 ± 0.7	19.0 ± 2.4	13.6 ± 0.8	18.0 ± 3.8
REP	16.4 ± 4.3	16.3 ± 1.4	19.3 ± 2.7	13.3 ± 1.7	18.0 ± 3.1
B ₁₂₀					
VEG	11.6 ± 2.3	16.0 ± 1.0	19.1 ± 2.5	12.7 ± 1.2	17.9 ± 3.3
REP	15.9 ± 5.1	15.9 ± 0.9	18.9 ± 2.3	13.0 ± 1.6	18.3 ± 3.0
R ₁₂₀					
VEG	11.8 ± 2.6	16.0 ± 1.0	19.1 ± 2.4	13.6 ± 0.8	17.8 ± 3.4
REP	15.9 ± 6.4	16.2 ± 1.4	19.5 ± 2.3	13.4 ± 1.2	18.3 ± 3.4

Table 3. Estimated phytochrome photoequilibria (PPE; P_{FR}/P_{R+FR}) and color fidelity index (CFI; R_f) of each supplemental lighting (SL) treatment. PPEs were calculated according to Sager et al. (1988), and CFI values were calculated according to supplemental materials provided by IES (2018). SL treatments consisted of either 460 W high-pressure sodium fixtures (HPS₁₂₀; LR48877; P.L. Light Systems, Beamsville, ON, Canada), 631 W light-emitting diode (LED) fixtures ($B_{20}G_{50}R_{45}FR_5$; VYPR 2p; Fluence, Austin, TX, USA), 325 W LED fixtures ($B_{20}R_{85}FR_{15}$; LumiGrow Pro 325; LumiGrow, Emeryville, CA, USA), 600 W LED fixtures ($B_{30}G_{25}R_{65}$; LX601G, Heliospectra, Göteborg, Sweden), a combination of 72 W LED fixtures (B_{120} ; HortiLED MULTI, P.L. Light Systems) and 625 W LED fixtures (R_{120} ; LumiGrow Pro 650E, LumiGrow), or 625 W LED fixtures (LumiGrow Pro 650E; LumiGrow).

	SL Treatment					
	HPS ₁₂₀	$B_{20}G_{50}R_{45}FR_5$	$B_{20}R_{85}FR_{15}$	$B_{30}G_{25}R_{65}$	B ₁₂₀	R ₁₂₀
Estimated PPE CFI (R _f)	0.85 44	0.85 80	0.84 0	0.87 55	0.50 <0	0.89 33

Plants were irrigated as needed with MSU Orchid RO Special (13 N–1.3 P–12.5 K watersoluble fertilizer containing (mg·L⁻¹) 125 nitrogen, 13 phosphorus, 121 potassium, 76 calcium, 19 magnesium, 1.7 iron, 0.4 copper and zinc, 0.9 manganese, 0.2 boron, and 0.2 molybdenum (GreenCare Fertilizers Inc.)), blended with reverse-osmosis water.



Figure 1. Emission spectra of supplemental lighting (SL) fixtures utilized throughout the study. SL treatments consisted of either 460 W high-pressure sodium fixtures (HPS₁₂₀; LR48877; P.L. Light Systems, Beamsville, ON, Canada), 631 W light-emitting diode (LED) fixtures ($B_{20}G_{50}R_{45}FR_5$; VYPR 2p; Fluence, Austin, TX, USA), 325 W LED fixtures ($B_{20}R_{85}FR_{15}$; LumiGrow Pro 325; LumiGrow, Emeryville, CA, USA), 600 W LED fixtures ($B_{30}G_{25}R_{65}$; LX601G, Heliospectra, Göteborg, Sweden), a combination of 72 W LED fixtures (B_{120} ; HortiLED MULTI, P.L. Light Systems) and 625 W LED fixtures (R_{120} ; LumiGrow Pro 650E, LumiGrow), or 625 W LED fixtures (LumiGrow Pro 650E; LumiGrow).

2.2. Data Collection and Analysis

Plants were monitored daily for the presence of the first visible flower bud (VB) and first open flower (OF). On the date of harvest (\geq 50 cm tall and three OFs for godetia; \geq 50 cm tall and inflorescence 50% open for snapdragon; \geq 45 cm tall and inflorescence 50% open for stock), stem length from the substrate surface to the tallest point of the inflorescence and caliper at the thickest point of the stem (recorded with a digital caliper (3-inch carbon fiber digital caliper, General Tools & Instruments, LLC, New York, NY, USA)) were recorded for all plants. Additionally, the total number of initiated inflorescences and branch number were recorded for snapdragon. A colorimeter (CR-20 Color Reader; Konica Minolta Sensing, Inc., Chiyoda, Tokyo, Japan) was utilized to measure flower petal color on three petals of each plant. Godetia flower color measurements were taken on the pink portion of the flower petal interiors. Data were analyzed using the SAS (version 9.4; SAS Institute, Cary, NC, USA) mixed-model procedure (PROC MIXED) for analysis of variance (ANOVA), and means were separated by Tukey's honest significant difference (HSD) test at $p \leq 0.05$. SAS general linear models procedure (PROC GLM) was used to fit regressions. Godetia and stock data were pooled across replications because of low harvestable stem yield and undetected single-flowering phenotypes being removed after transplant, respectively.

3. Results

3.1. Time to Visible Flower Bud

Time to VB (TVB) of godetia was influenced, albeit slightly, by the SL spectrum. TVB was the fastest for plants grown under HPS fixtures (52 d) and the slowest for plants grown under R_{120} SL (56 d). TVB was similar under all other treatments (\approx 53 d; Figure 2A). Snapdragon grown under $B_{20}R_{85}FR_{15}$, B_{120} , and HPS₁₂₀ SL reached VB the fastest (45–47 d), whereas TVB was delayed by up to 10 and 4 d under R_{120} SL during reps. 1 and 2, respectively (Figure 3A,B). TVB was delayed by 2–4 d and 1–3 d under $B_{20}G_{50}R_{45}FR_5$ and $B_{30}G_{25}R_{65}$ SL, respectively, compared to the fastest treatments. TVB of stock was the fastest for plants grown under B_{120}

SL (36 d). TVB was delayed by \approx 2, 3, 3, and 3 d when grown under B₂₀R₈₅FR₁₅, B₂₀G₅₀R₄₅FR₅,

 $B_{30}G_{25}R_{65}$, and HPS_{120} SL, respectively, compared to B_{120} SL. TVB was delayed by 9 d for



Figure 2. (**A**,**B**) time to visible flower bud, (**C**,**D**) time to open flower, (**E**,**F**) time to harvest, and (**G**,**H**) stem length at harvest of godetia 'Grace Rose Pink' and stock 'Iron Rose' in response to SL spectrum, pooled over two replications. SL treatments consisted of either 460 W HPS fixtures (HPS120; LR48877; P.L. Light Systems, Beamsville, ON, Canada), 631 W LED fixtures (B20G50R45FR5; VYPR 2p; Fluence, Austin, TX, USA), 325 W LED fixtures (B20R85FR15; LumiGrow Pro 325; LumiGrow, Emeryville, CA, USA), 600 W LED fixtures (B30G25R65; LX601G, Heliospectra, Göteborg, Sweden), a combination of 72 W LED fixtures (B120; HortiLED MULTI, P.L. Light Systems) and 625 W LED fixtures (R120; LumiGrow Pro 650E, LumiGrow), or 625 W LED fixtures (LumiGrow Pro 650E; LumiGrow). Means not followed by the same letter are significantly different by Tukey–Kramer honestly significant difference (HSD) test at $p \le 0.05$. Bars represent the mean and error bars indicate standard error.



Figure 3. (**A**,**B**) Time to visible flower bud, (**C**,**D**) time to open flower, (**E**,**F**) time to harvest, and (**G**,**H**) stem length at harvest of snapdragon 'Potomac Royal' in response to SL spectrum over two replications. SL treatments consisted of either 460 W HPS fixtures (HPS₁₂₀; LR48877; P.L. Light Systems, Beamsville, ON, Canada), 631 W LED fixtures (B₂₀G₅₀R₄₅FR₅; VYPR 2p; Fluence, Austin, TX, USA), 325 W LED fixtures (B₂₀R₈₅FR₁₅; LumiGrow Pro 325; LumiGrow, Emeryville, CA, USA), 600 W LED fixtures (B₃₀G₂₅R₆₅; LX601G, Heliospectra, Göteborg, Sweden), a combination of 72 W LED fixtures (B₁₂₀; HortiLED MULTI, P.L. Light Systems) and 625 W LED fixtures (R₁₂₀; LumiGrow Pro 650E, LumiGrow), or 625 W LED fixtures (LumiGrow Pro 650E; LumiGrow). Means not followed by the same letter are significantly different by Tukey–Kramer honestly significant difference (HSD) test at $p \leq 0.05$. Bars represent the mean and error bars indicate standard error.

3.2. Time to Open Flower

Godetia time to OF (TOF) was the fastest for plants grown under HPS SL (79 d) and the slowest for plants grown under B_{120} and R_{120} SL (88 and 91 d, respectively). TOF

was similar for all other SL treatments (84–86 d; Figure 2C). Snapdragon reached OF the fastest when grown under $B_{20}R_{85}FR_{15}$ and B_{120} SL during rep. 1 (65 and 66 d, respectively; Figure 3C,D). However, HPS₁₂₀ and $B_{20}R_{85}FR_{15}$ SL hastened flowering the most during rep. 2 (68 and 70 d, respectively). TOF was consistently delayed under R_{120} SL compared to the other treatments by up to 19 and 8 d during reps. 1 and 2, respectively. Flowering was slightly delayed under $B_{20}G_{50}R_{45}FR_5$ and $B_{30}G_{25}R_{65}$ SL compared to the fastest treatments during both reps, although by not as much as R_{120} SL. TOF was hastened for stock when grown under B_{120} and $B_{20}R_{85}FR_{15}$ SL (53 and 54 d, respectively). Flowering was delayed by 2–3 d for plants grown under $B_{30}G_{25}R_{65}$, HPS₁₂₀, and $B_{20}G_{50}R_{45}FR_5$ SL. Similar to TVB, TOF was delayed by 9 d when grown under R_{120} SL (Figure 2D).

3.3. Time to Harvest

Time to harvest (TTH) of godetia was the fastest under HPS₁₂₀ SL and the slowest under R₁₂₀ SL (80 and 94 d, respectively). TTH was 85 d for plants grown under B₂₀G₅₀R₄₅FR₅, B₂₀R₈₅FR₁₅, and B₃₀G₂₅R₆₅ SL, and 90 d for plants grown under B₁₂₀ SL (Figure 2E). TTH of snapdragon was hastened when grown under B₂₀R₈₅FR₁₅ and B₁₂₀ SL during rep. 1 (67 and 69 d, respectively; Figure 3E), while TTH was fastest under HPS₁₂₀ and B₂₀R₈₅FR₁₅ SL during rep. 2 (69 and 72 d, respectively; Figure 3F). TTH was slightly delayed when grown under B₂₀G₅₀R₄₅FR₅ and B₃₀G₂₅R₆₅ SL compared to the quickest treatments (4–5 d), while R₁₂₀ SL delayed harvest by up to 18 and 9 d during reps. 1 and 2, respectively. TTH of stock was the fastest when grown under B₁₂₀ and B₂₀R₈₅FR₁₅ SL (54 and 55 d, respectively). Flowering was delayed by \approx 2 d for plants grown under B₃₀G₂₅R₆₅, HPS₁₂₀, and B₂₀G₅₀R₄₅FR₅ SL. TTH was delayed by 10 d when grown under R₁₂₀ SL compared to B₁₂₀ SL (Figure 2F).

3.4. Cut Flower Morphology at Harvest

Godetia cut flower stems were the longest when grown under R₁₂₀ SL and the shortest when grown under B_{120} , $B_{20}G_{50}R_{45}FR_5$, and HPS_{120} SL (124 and 109–113 cm, respectively; Figure 2G). Godetia stem caliper was not influenced by SL treatment (p = 0.79). Snapdragon stems were the shortest when grown under B_{120} SL, regardless of rep. (Figure 3G,H). Plants were ≈ 13 or 24 cm longer when grown under R₁₂₀ SL during reps. 1 and 2, respectively. During rep. 1, stems grown under $B_{20}R_{85}FR_{15}$ SL were comparable in length to those grown under B₁₂₀ SL, although during rep. 2 they were approximately 10 cm longer. Similarly, under $B_{30}G_{25}R_{65}$ SL, stems were of similar length to those under B_{120} SL during rep. 1, while these stems were \approx 17 cm longer during rep. 2. Stems were of similar thickness regardless of SL treatment (11.6 to 12.9 mm). Moreover, snapdragon grown under B₂₀R₈₅FR₁₅, B₁₂₀, and HPS₁₂₀ SL had the fewest branches at harvest (52–55 branches), while plants grown under R_{120} SL produced stems with 8–11 more branches. The broad-spectrum LED fixtures produced stems with roughly five fewer branches than the R_{120} SL and up to six more branches than the other treatments. Snapdragon grown under R_{120} , B_{120} , and HPS_{120} SL had the fewest inflorescences at harvest, while $B_{20}R_{85}FR_{15}$ SL produced stems with \approx 5 more inflorescences. B₃₀G₂₅R₆₅ and B₂₀G₅₀R₄₅FR₅ SL yielded stems with 1–2 fewer inflorescences than B₂₀R₈₅FR₁₅ SL.

Stock stem length at harvest was commercially, but not statistically, similar between all treatments. B_{120} and R_{120} SL produced the longest stems (53 to 54 cm), while $B_{30}G_{25}R_{65}$ and $B_{20}R_{85}FR_{15}$ SL produced shorter cut flowers with an average stem length of \approx 50 cm (Figure 2H). Stock stem caliper was similar for all treatments except B_{120} , which produced stems up to 1.8 mm thinner than the other treatments.

3.5. Flower Petal Coloration at Harvest

Godetia and stock flower petal coloration was not influenced by any SL treatment (Table 4). Snapdragon petal coloration was not commercially different between treatments.

Table 4. Adjusted hue angle (h°), chroma (C), and Hunter CIELAB (L*, a*, b*) values at harvest for godetia, snapdragon, and stock grown under six different supplemental lighting (SL) treatments. SL treatments consisted of either 460 W high-pressure sodium fixtures (HPS₁₂₀; LR48877; P.L. Light Systems, Beamsville, ON, Canada), 631 W light-emitting diode (LED) fixtures (B₂₀G₅₀R₄₅FR₅; VYPR 2p; Fluence, Austin, TX, USA), 325 W LED fixtures (B₂₀R₈₅FR₁₅; LumiGrow Pro 325; LumiGrow, Emeryville, CA, USA), 600 W LED fixtures (B₃₀G₂₅R₆₅; LX601G, Heliospectra, Göteborg, Sweden), a combination of 72 W LED fixtures (B₁₂₀; HortiLED MULTI, P.L. Light Systems) and 625 W LED fixtures (R₁₂₀; LumiGrow Pro 650E, LumiGrow), or 625 W LED fixtures (LumiGrow Pro 650E; LumiGrow). Letters indicate mean separations across treatments using Tukey–Kramer honestly significant difference (HSD) test at $p \leq 0.05$.

	SL Treatment							
Parameter	HPS ₁₂₀	$B_{20}G_{50}R_{45}FR_5$	B ₂₀ R ₈₅ FR ₁₅	$B_{30}G_{25}R_{65}$	B ₁₂₀	R ₁₂₀		
		Godetia 'Grace Rose Pink'						
h°	355.2 ^{NS}	354.9	354.5	354.3	353.6	355.2		
С	48.5 ^{NS}	49.1	47.7	49.4	47.4	50.1		
L*	43.2 ^{NS}	42.8	44.8	42.5	44.5	43.8		
a*	48.3 ^{NS}	48.9	47.5	49.1	47.1	49.9		
b*	-3.9 NS	-4.2	-4.5	-4.4	-5.2	-3.7		
		Snapdragon 'Potomac Royal'						
h°	355.1 C	356.8 A	357.2 A	356.3 AB	356.9 A	355.7 BC		
С	34.4 A	34.2 AB	33.0 BC	33.7 ABC	32.7 C	34.3 ABC		
L*	19.4 B	19.7 B	19.6 B	20.2 B	19.8 B	21.7 A		
a*	34.5 A	34.2 AB	33.0 BC	33.7 ABC	32.7 C	34.2 ABC		
b*	-2.7 D	-1.8 BC	-0.9 A	-1.8 B	-1.3 AB	-2.6 CD		
	Stock 'Iron Rose'							
h°	336.8	337.3	337.0	336.9	337.5	337.0		
С	49.6	49.2	49.5	49.1	49.9	49.8		
L*	33.9	33.9	34.1	33.6	34.1	33.3		
a*	45.6	45.3	45.6	45.1	46.0	45.8		
b*	-19.5	-19.0	-19.3	-19.3	-19.1	-19.5		

Means not followed by the same letter are significantly different by by Tukey–Kramer honestly significant difference (HSD) test at $p \le 0.05$.

4. Discussion

With a variety of commercially available SL fixtures on the market, it is important to understand the influence that the supplemental radiation quality can have on the growth and development of cut flowers. We found that development time, in addition to cut flower morphology, varied between the spectra that were studied. Generally, TVB, TOF, and TTH were the slowest for plants grown under R_{120} SL, regardless of variety. However, the varieties studied exhibited different developmental responses to the remaining SL spectra. Godetia consistently developed the fastest under HPS₁₂₀ SL. Stock developed the fastest when grown under B_{120} and $B_{20}R_{85}FR_{15}$ SL, while snapdragon consistently developed the fastest when grown under $B_{20}R_{85}FR_{15}$, B_{120} , and HPS₁₂₀ SL.

While R radiation alone is sufficient to inhibit flowering in most short-day plants, many LDPs require R and FR radiation to induce flowering, particularly when the DLI is low (e.g., <8 mol·m⁻²·d⁻¹). Craig and Runkle [21] reported that flowering of snapdragon 'Liberty Classic Cherry' was delayed by up to 14 d when grown under a 4 h NI provided by $\approx 1.5 \ \mu mol \cdot m^{-2} \cdot s^{-1}$ of R radiation (PPE = 0.89) compared to the same NI provided by both R and FR radiation (PPE = 0.72). This phenomenon may have contributed to the developmental delay seen under R₁₂₀ SL across all genera (Figures 4 and 5), which had an equivalent PPE of 0.89. While FR radiation from solar radiation was available for plants under each treatment, SL emitting a moderate flux of FR radiation reduced the estimated PPE and appeared to hasten plant development. The same is true of the B₁₂₀ SL treatment, which reduced the estimated PPE by 0.39 compared to the R₁₂₀ treatment.



Figure 4. Influence of estimated phytochrome photoequilibrium (PFR/PR+PFR) of supplemental lighting treatments (**A**,**B**) on time to visible bud, (**C**,**D**) time to open flower, (**E**,**F**) time to harvest, and (**G**,**H**) stem length at harvest of godetia 'Grace Rose Pink' and stock 'Iron Rose'. Black symbols represent means; error bars represent standard error. \mathbb{R}^2 values are presented; *** indicate model significance at *p* < 0.00001. Coefficients are presented in Table 5.

The effect of B_{120} SL on development time varied between the LDPs studied. While stock and snapdragon experienced hastened development when grown under B_{120} SL, development of godetia slowed when grown under B_{120} SL compared to most of the other treatments, indicating that this response may be genus specific. This is supported by Hori et al. [44], who reported that baby's breath 'Bristol Fairy' (*Gypsophila paniculata*) did not flower when grown under a 12 h DE provided by 20–30 µmol·m⁻²·s⁻¹ of B radiation for 18 weeks. However, flowering occurred after \approx 75 or 98 d when plants were grown under 9 μ mol·m⁻²·s⁻¹ of incandescent lighting or 20–30 μ mol·m⁻²·s⁻¹ of FR radiation for the same duration, respectively [44]. TVB, TOF, and TTH of stock and snapdragon was delayed as the estimated PPE increased from 0.50 (B₁₂₀) to 0.89 (R₁₂₀; Figures 4 and 5), indicating that the developmental delay between SL treatments could be at least partly due to increased phytochrome activity. This is in agreement with Craig and Runkle [23], who reported that TOF of the LDPs petunia 'Easy Wave White' (*Petunia* ×*hybrida*) and snapdragon 'Liberty Classic Cherry' was delayed by up to ≈6 d and 12 d, respectively, as the estimated PPE of NI lighting increased from 0.46 to 0.89.

Table 5. Regression equations and R² for time to visible bud, time to open flower, time to harvest, and stem length at harvest of godetia 'Grace Rose Pink', stock 'Iron Rose', and snapdragon 'Potomac Royal' in response to the estimated phytochrome photoequilibrium of each supplemental lighting treatment. ** and *** indicate model significance at p < 0.001 and p < 0.0001, respectively. All models are in the form of: $f = y0 + a * PPE + b * PPE^2$.

Parameter	y0	Α	b	R ²			
	Godetia 'Grace Rose Pink'						
Time to visible bud (d)	136.93 ^z	-265.74	196.25	0.206 ***			
Time to open flower (d)	324.77	-746.35	543.98	0.371 ***			
Time to harvest (d)	355.71	-841.28	613.57	0.400 ***			
Stem length at harvest (cm)	352.60	-785.44	593.81	0.151 ***			
		Stock 'Iron	Rose'				
Time to visible bud (d)	163.75	-410.27	309.94	0.460 ***			
Time to open flower (d)	203.47	-485.22	366.37	0.503 ***			
Time to harvest (d)	212.48	-510.96	385.82	0.422 ***			
Stem length at harvest (cm)	104.79	-160.00	113.78	0.081 ***			
	Snapdragon 'Potomac Royal'						
Time to visible bud (d)							
Rep. 1	220.56	-563.92	423.74	0.621 ***			
Rep. 2	128.57	-259.47	193.97	0.334 ***			
Time to open flower (d)							
Rep. 1	429.16	-1166.49	874.18	0.782 ***			
Rep. 2	221.23	-469.85	344.12	0.380 ***			
Time to harvest (d)							
Rep. 1	410.51	-1097.96	822.36	0.681 ***			
Rep. 2	232.81	-500.12	366.46	0.398 ***			
Stem length at harvest (cm)							
Rep. 1	362.45	-760.81	586.91	0.268 ***			
Rep. 2	264.58	-424.62	333.72	0.112 **			

^z Coefficients for model equations were used to generate Figures 4 and 5.

Both $B_{20}G_{50}R_{45}FR_5$ and $B_{30}G_{25}R_{65}$ SL slightly delayed development compared to the fastest treatments, but not as significantly as R_{120} SL. This delay could be attributed to the minimal emission of FR radiation in the former treatment and the lack of FR radiation in the latter treatment, which resulted in higher estimated PPEs (0.85 to 0.87), although not as high as R_{120} SL (0.89). Moreover, this delay may have lasted longer if these spectra did not contain B and G radiation, as both wavebands can serve as long-day signals when applied at moderate intensities.

SL quality also influenced cut flower morphology. Stem lengths at harvest were generally the shortest under B_{120} SL regardless of genus and increased with the estimated PPE (Figures 4 and 5). Many floriculture crops exhibit a compact growth habit when grown under B radiation. For instance, Zou [45] found that geranium 'Calliope Dark Red' plants grown with 100% B radiation for 24 h·d⁻¹ was up to 6.7 cm wider in comparison to those grown with 100% R radiation for 24 h·d⁻¹. Moreover, baby's breath 'Bristol Fairy' grown under a 24 h photoperiod created with 16 h of DE lighting providing 20–30 µmol·m⁻²·s⁻¹ of B radiation was ≈43 cm shorter than those grown under the same intensity and duration provided by 100% FR radiation [44]. This compaction could be at least partly regulated by

phytochrome activity. Kong et al. [46] found that continuous exposure of 100 μ mol·m⁻²·s⁻¹ of B radiation for 14–20 d promoted stem elongation of several bedding plants compared to the same intensity and duration of R radiation. However, when \approx 90 μ mol·m⁻²·s⁻¹ of B radiation was applied with an additional flux of \approx 10 μ mol·m⁻²·s⁻¹ of R radiation, plants were more compact than any other treatment. The authors concluded that this response could be due to reduced phytochrome activity under sole-source B radiation (PPE = 0.49), promoting stem elongation, compared to a combination of B and R radiation (PPE = 0.74). Considering that plants in the present study were grown in greenhouses with solar and supplemental radiation, the actual PPE under B₁₂₀ SL would likely be >0.50 because of the presence of other wavebands, potentially contributing to our similar findings. However, stock cut flowers were the longest when grown under B₁₂₀ and R₁₂₀ SL and the shortest when grown under B₃₀G₂₅R₆₅ and B₂₀R₈₅FR₁₅ SL, though differences were minimal.



Figure 5. Influence of estimated phytochrome photoequilibrium (PFR/PR+PFR) of supplemental lighting treatments on (**A**,**B**) time to visible bud, (**C**,**D**) time to open flower, (**E**,**F**) time to harvest, and (**G**,**H**) stem length at harvest of snapdragon 'Potomac Royal' during replications 1 and 2. Black symbols represent means; error bars represent standard error. R² values are presented; ** and *** indicate model significance at *p* < 0.001 and *p* < 0.0001, respectively. Coefficients are presented in Table 5.

This further supports the argument that B-mediated stem elongation is a genus-specific response. Another instance of B-mediated stem elongation was published by Zou [45],

who found that marigold 'P-4' (*Tagetes erecta*) grown under sole-source lighting providing 180 μ mol·m⁻²·s⁻¹ of B radiation for 12 h·d⁻¹ was up to 54% taller than those grown under 180 μ mol·m⁻²·s⁻¹ of R radiation for the same duration. It was also found that petunia and dianthus seedlings grown under SL emitting 19% B radiation for 16 h·d⁻¹ were 59% and 3% taller, respectively, than those grown under SL emitting 6% B radiation [47]. However, the former SL treatment included 5% G radiation, which may have antagonized B-mediated stem compaction compared to the latter treatment, which did not contain G radiation.

Additionally, snapdragon grown under $B_{20}R_{85}FR_{15}$ SL (estimated PPE = 0.84) had \approx 5 more inflorescences at harvest compared to those grown under R_{120} SL (estimated PPE = 0.89). This contrasts with Craig and Runkle [21], who found that snapdragon 'Liberty Classic Cherry' had up to eight more VBs when grown under 100% R NI lighting (estimated PPE = 0.89) compared to other NI treatments creating an estimated PPE of 0.16 to 0.85.

The present study demonstrates the influence that SL quality can have on crop growth and development. However, these effects cannot be relied on year-round as a means of crop steering and growth regulation, as the effects of SL quality on crop growth and developmental responses are the strongest when the solar DLI is low [48,49]. For instance, when the quotient of B radiation provided by SL increased from 0% to 30% when SL provided 45–70% of the total DLI (ranging from 2.1–8.4 mol·m⁻²·d⁻¹), stem elongation of celosia 'Fresh Look Gold' (Celosia argentea), snapdragon 'Rocket Pink', and vinca 'Titan Punch' (Catharanthus roseus) was suppressed by $\approx 20\%$, $\approx 10\%$, and $\approx 30\%$, respectively [48,50]. In a separate study, where the DLI was consistently > 6.7 mol \cdot m⁻²·d⁻¹ and SL only provided 20–40% of the total DLI, there was no commercial effect on seedling stem elongation as the quotient of B radiation provided by SL increased from 10% to 45% [10,48]. Moreover, Hernández and Kubota [51] reported no statistical morphological differences between greenhouse-grown tomato seedlings grown with SL of varying spectra and a DLI of either 8.9 or $19.4 \text{ mol} \cdot \text{m}^{-2} \cdot \text{d}^{-1}$. These findings indicate that while SL spectrum may be less influential as the DLI increases, particularly > 7 mol·m⁻²·d⁻¹, it can have noticeable effects on crop growth and development when the solar DLI is below this threshold.

While parameters including TOF, TTH, and finished stem quality must be considered when selecting a spectrum for a SL strategy, human work suitability must also be considered. In the present study, $B_{20}R_{85}FR_{15}$, B_{120} , and HPS_{120} SL consistently hastened plant development and yielded cut flowers with moderate stem lengths. Conversely, R_{120} SL delayed TTH and produced longer cut flowers. While a given spectrum may elicit desirable crop responses, it may create a challenging work environment for humans by making it more difficult to diagnose cultural issues, including nutrient deficiencies and pest prevalence on plant tissue [52]. This may be particularly true when the solar DLI is low, and SL contributes more to the total DLI than solar radiation.

The color fidelity index (CFI; R_f) is an independent, unbiased indication of how well natural colors can be perceived by the human eye under a particular light source [34]. The CFI exists on a scale of 0 to 100, where values closer to 100 indicate that the colors perceived under a given light source are truer to nature [34]. The CFI values of each SL treatment were calculated with each source's spectral power distribution according to supplemental materials provided by IES [52] and can be found in Table 3. While B₂₀R₈₅FR₁₅ SL generally hastened TTH and produced stems with moderate lengths, it created an environment with a lower CFI than HPS_{120} SL or either broad-spectrum fixture, meaning that human visibility capacity would be impaired under that spectrum. However, the effects of $B_{20}R_{85}FR_{15}$ SL's low CFI were the strongest during the early morning and evening, while solar radiation was limiting. During the day, the higher fraction of solar radiation subjectively allowed for sufficient human visibility. Additionally, both B_{120} ($R_f < 0$) and R_{120} SL ($R_f = 33$) created environments that were inadequate for human visibility and sufficient crop supervision (Table 3). Similar to $B_{20}R_{85}FR_{15}$ SL, the impact on visibility by these treatments was the strongest when solar radiation was limiting; however, visibility was still noticeably impaired during the day compared to any other treatments.

SL fixtures also vary in their capability to convert electrical power to photons. Photon efficacy is defined as the number of moles of photons generated per energy input, typically expressed as μ mol·J⁻¹ [8,53]. Currently, LED fixtures can have a photon efficacy of up to 3 μ mol·J⁻¹, trumping the photon efficacy of HPS fixtures by approximately 60%. This is partly because a substantial amount of energy consumed by HPS fixtures is re-emitted as heat, whereas LED fixtures typically function at a lower temperature. This can have a significant impact on a greenhouse operation's overall energy expenditure. While more energy must be used to heat a greenhouse when using LED fixtures compared to HPS fixtures, the net energy expenditure, and associated energy costs, can be 10–25% lower than greenhouses utilizing HPS fixtures [53].

Rate of development, finished stem quality, crop visibility, and photon efficacy must be considered when selecting a SL spectrum for one's growing operation. Based on our findings, we recommend utilizing an LED fixture that provides a light ratio similar to $B_{20}R_{85}FR_{15}$ SL or broad-spectrum light; both elicited desirable crop responses with minimal tradeoffs, while allowing for sufficient human visibility. Although crops grown under HPS₁₂₀ SL performed similarly, we recommend utilizing LEDs as they most likely offer higher photon efficacy and the potential for long-term energy and monetary savings.

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