Spray application factors and plant growth regulator performance: IV. Dose response relationships

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Abstract. Effect of carrier volume (range 119 to 668 L·ha⁻¹) on dose response relationships of daminozide and GA₃ was investigated using bean (Phaseolus vulgaris L.) seedlings as a model system. Carrier volume was varied by altering nozzle travel speed thereby maintaining constant flow rate and droplet size. Response was indexed by inhibition (daminozide) or stimulation (GA₃) of elongation of first plus second internodes above primary leaves 14 days after spray application. Increasing dose by increasing concentration and/or increasing carrier volume at constant concentration increased response. For a given dose retained, response to daminozide was related positively to carrier volume, while GA₃ response was not affected. Chemical names used: butanedioic acid mono(2,2-dimethylhydrazide) (daminozide); gibberellic acid (GA₃).

To investigate the effect of carrier volume on performance of plant growth regulators in a more critical manner, we selected a model system using bean (Phaseolus vulgaris) seedlings and the growth regulators daminozide and GA₃. These compounds were chosen, because 1) they represent growth regulators differing in polarity, 2) both induce rapid and marked effects on shoot elongation that can be quantified readily, and 3) information on droplet size and volume effects is available under controlled conditions (Knoche et al., 1998; Knoche and Bukovac, 1999 and 2000).

Materials and Methods

Plant material. Bean seed of two cultivars, determine ‘Green Ruler’ (GR) for daminozide assay and indeterminate ‘Black Seeded Blue Lake’ (BSBL) for GA₃ assay, were sown (three or four seeds per pot) in peat pots (0.58-L volume) filled with a commercial growing medium (PROMIX BW; Premier Brands, New Rochelle, N.Y.). Following emergence, plants were selected for uniformity and freedom from defects and thinned to one plant per pot. Plants were watered daily. Fertilizer was provided either initially with the growing medium (14N–4.2P–11.6K Osmocote at 8 g·L⁻¹; Scotts-Sierra Horticultural Prod. Co., Marysville, Ohio) or at weekly intervals by fertigation with Peters 20N–8.8P–16.6K water-soluble fertilizer (Scotts-Sierra Horticultural Prod. Co.) at an N concentration equivalent to 50 mg·L⁻¹. Growing conditions were a growth chamber set at 16-h day/8-h night temperatures of 25/20 ± 2 °C and 45/62 ± 5% relative humidity with a 16-h photoperiod provided by cool-white fluorescent lamps (F72T12 CW-1500; General Electric Co., Cleveland, Ohio) at 200 µmol·m⁻²·s⁻¹ photosynthetic active radiation, as measured with a quantum–radiometer–photometer (LI-185; LI-COR, Lincoln, Nebr.), at plant level and supplemented by 1% red light (tungsten filament bulbs). Plants were maintained under these conditions for the entire experimental period except when removed (≈2 h) for treatment 7 or 8 d after seeding.

Growth regulators. Spray solutions were prepared with deionized water using daminozide (>99% pure, technical grade; Uniroyal Chemical Corp., Middlebury, Conn., or analytical grade; Sigma Chem. Co., St. Louis, Mo.) and gibberellic acid (GA₃, content >90%; Sigma Chemical Co.). GA₃ concentrations were based on a GA₃ content of 100%.

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SPRAY APPLICATION AND BIOLOGICAL RESPONSE. Effect of carrier volume on daminozide inhibition and GA₃ promotion of internode elongation was investigated following spray application using an experimental track sprayer. Concentrations of spray solutions for daminozide and GA₃ were 150, 300, 600, 1200, and 2400 mg·L⁻¹ and 2.5, 5, 10, 20, and 40 mg·L⁻¹, respectively. The pH of daminozide solutions averaged 3.6 (range 3.5 to 3.9; water control 5.5), while pH of GA₃ solutions decreased from 4.8 to 3.8 as concentration increased from 2.5 to 40 mg·L⁻¹. The track sprayer was equipped with a conventional flat fan spray nozzle (8002; Spraying Systems, Wheaton, Ill.) mounted on a variable speed transport carriage. Seedlings were positioned ≈46 cm below the nozzle with primary leaves orientated perpendicular to travel of the nozzle. The flow rate averaged 258 mL·min⁻¹ at 39.6 kPa measured at the spray nozzle. Specified carrier volumes (range 138 to 668 L·ha⁻¹) were achieved by varying travel speed (3.6 to 0.75 km·h⁻¹). Orifice size and flow rate were kept constant. Under these conditions, volume median diameter of spray droplets was $356 \pm 43 \mu m$ (SE, n = 3). Sprays were applied to primary bean leaves at the midpoint of the photoperiod ±1 h at 25 ± 3°C. Plants were selected further so that the length of the first internode was <1 cm. When necessary, leaves were supported from below (deviation from horizontal <20°) to ensure a similar and full plan leaf area exposure to the spray. Plants were returned immediately to the growth chamber after spraying, randomized, staked with bamboo when dry, and allowed to grow for 14 d. Unless otherwise specified, elongation of first plus second internodes was determined with 10 replications per concentration. Dose response relations established in previous studies may be linearized using the following transformations (Knoche et al., 1998). For daminozide, log internode length (cm) = –0.27 × log (concn, g·L⁻¹) + 0.79, $r^2 = 0.940^*$; for GA₃, internode length (cm) = 19.0 × log (concn, mg·L⁻¹) + 18.8, $r^2 = 0.986^*$. Data for the highest GA₃ concentration were excluded from this analysis, since GA₃ internode elongation response was saturated at doses exceeding 7.5 µg/leaf (see Fig. 2B in Knoche et al., 1998).

SPRAY RETENTION, COVERAGE, AND SURFACE TENSION. Retention was measured gravimetrically for each pass of the nozzle. Primary leaves of comparable plants were excised, weighed, positioned in the spray path at the same height and orientation as for the record plants, sprayed, and then reweighed. Leaf area was determined using a leaf area meter (model LI-3000; LI-COR). A maximum retention value of $6.7 \mu L·cm⁻²$ (equivalent to 665 L·ha⁻¹) was calculated when spraying plants just short of runoff.

Spray distribution and leaf coverage were assessed by spraying leaves with a 2% nigrosin dye solution using the same sprayer settings. Leaves were allowed to dry, excised, positioned between layers of botanical blotting paper, pressed, and stored in the

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**Table 1. Documented responses of growth regulator performance when applied in reduced carrier volumes in tree fruit species.**

| Growth regulator | Species | Carrier vol range (L·ha⁻¹) | Response index | Effect on performance⁴ | Reference |
|------------------|---------|---------------------------|---------------|------------------------|----------|
| Accel⁵           | Apple   | 500–2000                  | Fruit size, yield | No effect              | Hull et al. (1995) |
| Daminozide       | Apple   | 93–1870                   | Fruit drop     | Decreased              | Rogers and Krestensen (1973) |
|                   | Sour cherry | 94–2350                 | Fruit removal force | No effect              | Bukovac (1981) |
| Ethephon         | Sweet cherry | 118–2350                | Fruit removal force | Increased              | Bukovac (1981) |
|                   | Walnut   | 935–2804                 | Nut removal (%), hullability | Increased              | Olson et al. (1977) |
| Fenoprop⁶        | Apple   | 159–3972                  | Fruit drop     | No effect              | Bukovac, unpublished |
| GA₃              | Sour cherry | 234–1738                 | Flowering      | No effect              | Bukovac, unpublished |
| NAA              | Apple   | 230–2100                  | Thinning, fruit size | No effect              | Black et al. (1994) |

⁴Effect relative to response obtained from high volume application.
⁵Formulation of benzyladenine (1.8%) + GA₃(0.18%).
⁶Formulation of 2-(2,4,5-trichlorophenoxy) propionic acid.

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Fig. 1. (A) effect of nozzle travel speed on the spray volume retained per unit leaf area of primary leaves of bean seedlings (inset: spray retention vs. the inverse of travel speed) and (B) effect of carrier volume on actual percentage leaf area covered compared with theoretical coverage expected if spray impacted and remained and dried as discrete droplets.
Fig. 2. Representative images of spray deposition on primary leaves of bean at carrier volumes of (A) 653, (B) 483, (C) 258, and (D) 142 L·ha⁻¹ applied with a flat-fan nozzle. Corresponding leaf area covered was 34.4%, 31.3%, 21.0%, and 15.3%, respectively. Each image represents an original area sampled of 0.85 cm × 1.24 cm or 1.05 cm².

Fig. 3. Effect of spray concentration and carrier volume on (A) daminozide inhibition and (B) GA₃ promotion of internode elongation in bean seedlings.

Fig. 4. (A) daminozide inhibition and (B) GA₃ promotion of internode elongation of bean seedlings as a function of dose retained per unit leaf area. For regression equations see Table 2.
travel speed against spray retention yielded a significant linear relationship, Retention (µL·cm⁻²) = 7.2 × speed⁻¹ (h·km⁻¹) – 0.88, \( r^2 = 0.977^{***} \), indicating that retention efficiency was independent of the spray volume applied at the speeds and volumes used (Fig. 1A, inset). This may be expected, since 1) droplet size was maintained constant and that changing travel speed only altered the number of droplets impacting on the leaf surface, 2) surface tension of daminozide and GA₃ solutions were independent of concentration and not significantly different from water, and 3) there was no significant runoff of spray solution from the leaf surface in this experiment.

A positive relationship was obtained between leaf coverage and the volume applied (Fig. 1B). Increasing spray volume from 142 to 653 L·ha⁻¹ increased spray coverage from 15.3% to 36.7% (Fig. 1B and 2). However, the increase in coverage was less than expected if droplets impacted and spread independently (solid vs. dotted line, Fig. 1B). This observation suggested that an increasing number of droplets coalesced so that deposits overlapped as carrier volume was increased (see also Fig. 2).

Effects of daminozide on inhibition and GA₃ on promotion of internode elongation were related positively to the concentration of the active ingredient (a.i.) and/or carrier volume (Fig. 3A and B). Since spray retention has been measured as a function of nozzle travel speed, the amount of daminozide and GA₃ retained per unit leaf area can be calculated from retention data for any given concentration and carrier volume. Linearizing dose response relations by appropriate transformations noted earlier and plotting response vs. dose retained per unit leaf area yielded the relationships illustrated in Fig. 4 (for regression equations see Table 2). Since a constant concentration range was applied at different carrier volumes, the amounts of a.i. retained per unit leaf area differed among spray volumes. However, doses in the ranges of 1.5 to 2.8 µg·cm⁻² (daminozide) and 0.022 to 0.042 µg·cm⁻² (GA₃) were common to all carrier volumes (Fig. 4A and B), and for these ranges, internode elongation was calculated as a function of carrier volume.

Increasing carrier volume at constant daminozide dose decreased internode length (Fig. 5A). In contrast, GA₃ promotion of internode elongation was independent of carrier volume (Fig. 5B). Absence of a carrier volume effect on GA₃ performance was confirmed, when an approximately constant dose of GA₃ per unit leaf area was applied in various carrier volumes (Table 3). While GA₃ increased internode length compared to the control, there was no difference in response to a given dose delivered in different carrier volumes (Table 3). These data are consistent with

### Results and Discussion

Increasing nozzle travel speed decreased the volume of spray solution retained per unit leaf area (Fig. 1A). Plotting the inverse of travel speed against spray retention yielded a significant linear relationship, Retention (µL·cm⁻²) = 7.2 × speed⁻¹ (h·km⁻¹) – 0.88, \( r^2 = 0.977^{***} \), indicating that retention efficiency was independent of the spray volume applied at the speeds and volumes used (Fig. 1A, inset). This may be expected, since 1) droplet size was maintained constant and that changing travel speed only altered the number of droplets impacting on the leaf surface, 2) surface tension of daminozide and GA₃ solutions were independent of concentration and not significantly different from water, and 3) there was no significant runoff of spray solution from the leaf surface in this experiment.

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### Table 2. Regression equations for dose response relationships for daminozide or GA₃ and internode elongation in bean seedlings.

| Growth regulator | Carrier vol (L·ha⁻¹) | Regression coefficient a ±SE | Coefficient of determination \( r^2 \) |
|-----------------|----------------------|-------------------------------|---------------------------------|
| Daminozide      | 119                  | -0.11 ± 0.05                 | 0.605*                          |
|                 | 159                  | -0.40 ± 0.09                 | 0.873*                          |
|                 | 311                  | -0.72 ± 0.08                 | 0.968**                         |
|                 | 665                  | -0.33 ± 0.11                 | 0.621†                          |
| GA₃             | 138                  | 9.1 ± 0.7                     | 0.981**                         |
|                 | 213                  | 10.3 ± 1.1                    | 0.969**                         |
|                 | 378                  | 11.6 ± 0.9                    | 0.981**                         |
|                 | 668                  | 17.6 ± 1.6                    | 0.975**                         |

*Internode elongation response was indexed by length of first plus second internodes 14 d after treatment. Regression equations were of the form log internode length (cm) = a × log (dose, µg·cm⁻²) + b for daminozide and Internode length (cm) = a × log (dose, µg·cm⁻²) + b for GA₃.

**Nonsignificant or significant at \( P \leq 0.05 \) or 0.01, respectively.

Fig. 5. Predicted effects of carrier volume on (A) daminozide-inhibition and (B) GA₃ promotion of elongation of first plus second internodes of bean seedlings. Vertical bars represent 95% confidence limits. For details see text.
Table 3. Effect of carrier volume at constant dose of GA$_3$ on internode elongation in bean seedlings.

| Carrier vol (L·ha$^{-1}$) | Concen (mg·L$^{-1}$) | GA$_3$ retention (µg·cm$^{-2}$) | Internode length$^2$ (cm) |
|--------------------------|----------------------|-------------------------------|--------------------------|
|                          |                      |                              | First        | Second       | Third        |
| Control                  | ---                  | ---                           | 3.2 a        | 2.5 a        | 5.5 a        |
| 160                      | 40                   | 0.046                         | 4.0 a        | 5.0 b        | 12.6 b       |
| 269                      | 20                   | 0.053                         | 4.2 a        | 5.0 b        | 12.5 b       |
| 420                      | 10                   | 0.041                         | 3.7 a        | 5.5 b        | 13.2 b       |
| 730                      | 5                    | 0.047                         | 4.2 a        | 5.7 b        | 13.3 b       |

$^2$Mean separation within columns by Duncan’s multiple range test, $P \leq 0.05$.

earlier observations following droplet application of daminozide and GA$_3$ to primary bean leaves (Knoche et al., 1998). Since retention efficiency on the easy-to-wet bean leaf surface was independent of carrier volume (Fig. 1A), effects on the growth regulator induced growth responses were most likely related to a carrier volume effect on foliar uptake and/or subsequent transfer stages. In our earlier studies we observed higher translocation (percentage of applied) at higher carrier volumes (Knoche and Bukovac, 1999 and 2000). Unfortunately, the coverage range investigated was markedly narrower (0.3% to 10.6%) compared with the present study (15.3% to 36.7%). Also, in our earlier study (Knoche and Bukovac, 2000) we observed phytotoxicity at low carrier volumes/high concentrations of daminozide. Thus, we do not know if the relationship obtained earlier fully applies to our present study, where no toxicity was observed.

Care must be taken in extrapolating our findings to other species that differ significantly in wettability. For spray solutions containing surfactants, retention efficiency and solution concentration, i.e., carrier volume, may be interrelated (Anderson et al., 1983; Reichard, 1988; Thomas and Hall, 1979). For difficult-to-wet leaf surfaces (contact angles $>90^\circ$) retention efficiency is expected to increase with decreasing carrier volume and this would increase performance. Furthermore, canopy architecture of crops may differ dramatically from the model system employed in our study and altering spray application parameters may alter canopy penetration of spray droplets thereby affecting performance.

In conclusion, our data confirm that performance of growth regulators may be affected by carrier volume. Hence, for a given response, the dose of a compound applied may be a function of carrier volume. At present, the data base for adjusting doses for a carrier volume effect is very limited. Research in this area is critically needed not only to increase plant growth regulator performance, but also to increase spray efficiency and reduce nonessential chemical input into the environment.

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