Crop Injury from Sublethal Rates of Herbicide. III. Pepper

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Abstract. Sublethal rates of 2,4-D and dicamba were applied to pepper to evaluate the possible effects of single or multiple exposures to drift from these herbicides.Dicamba induced more foliar injury than did 2,4-D and reduced vigor more as herbicide rates increased. Postbloom applications reduced vigor less than earlier applications. Epinastic response was affected by stage of development at application and time after treatment. Postbloom applications did not affect yield, but dicamba and 2,4-D at early stages of development resulted in linear reduction of marketable and total yields as rates increased to 112 g·ha⁻¹. Reductions in plant vigor with increased rates were greater and foliar epinasty was more pronounced with two sequential applications of 2,4-D or dicamba than with single applications. Marketable yields were unaffected by single prebloom applications but declined linearly with two applications. Cull and total yields were not affected by the number of applications. With prebloom and bloom applications of 2,4-D, flower abscission increased and fruit set decreased as rate increased. Chemical names used: 3,6-dichloro-2-methoxybenzoic acid (dicamba); 2,4-dichlorophenoxy)acetic acid (2,4-D).

Dicamba and 2,4-D selectively control broadleaf weeds in grass crops. In sensitive plants, these auxin-like herbicides stimulate ethylene formation (Abeles, 1968). Phytotoxicity results from the accumulation of cyanide, a coproduct of ethylene synthesis (Tittle et al., 1990). In Florida, both 2,4-D and dicamba are registered for broadleaf weed control in corn (Zea mays L.), sorghum (Sorghum bicolor L. Moench), small grains, turf, pasture and range-land. Spray and vapor drift of auxinic herbicides can be damaging to nontarget broadleaf crops. The use of 2,4-D along roadways during windy conditions has resulted in damage to crops in adjacent fields (Gilreath, personal observation).

Spraying with sublethal concentrations is a commonly used technique to evaluate the possible effects of herbicidal drift. In assessing the effects of sublethal rates of dicamba on potato (Solanum tuberosum L.), foliar injury and stunting were reported with rates of 2.8 to 22.2 g·ha⁻¹, and total and marketable yields were reduced (Wall, 1994). Similar work on tomato (Lycopersicon esculentum Mill.) has been focused primarily on the effects of 2,4-D vapor drift, with fewer reports of spray drift effects on processing tomato. In field and greenhouse studies, foliar injury and abnor-mal fruit were reported with exposure to 2,4-D vapor (Carlin et al., 1971; Sherwood et al., 1970; Weigle et al., 1970). Incidence of injury was directly related to 2,4-D vapor concentrations in laboratory studies (Bennet, 1989; Breeze, 1988; Breeze and van Rensburg, 1991, 1992; Breeze and West, 1987). Yields of processing tomato were more adversely affected by sublethal rates of 2,4-D and dicamba when exposure occurred during flowering than after the fruits were set (Jordan and Romanowski, 1974).

Less information is available on the effects of sublethal concentrations of 2,4-D on pepper (Capsicum annuum L.) than on tomato, and no information is available on the effects of low rates of dicamba on pepper. Pepper plants developed foliar symptoms of phenoxy herbicide injury after exposure to road or field dust containing residues of an amine formulation of 2,4-D (Simons and Pariente, 1982). With Capsicum frutescens L., 104 g·ha⁻¹ 2,4-D induced foliar epinasty, stunting of shoots, and flower bud abscission, and reduced yield, whereas 2 g·ha⁻¹ increased yield (Hemphill and Montgomery, 1981). Orsenigo (1964) observed foliar symptoms, abscission of flowers and small fruit, and abnormal fruit development following exposure of pepper to phenoxy herbicides.

We determined effects on pepper growth and yield of sublethal rates of dicamba and 2,4-D applied as single and multiple applications at various concentrations and at different stages of development. This study expands on that of Hemphill and Montgomery (1981) by examining the more commonly cultivated C. annuum, evaluating the effects of multiple exposures, comparing the effects of two herbicides, and using the polyethylene-mulch production system commonly used for pepper production.

Materials and Methods

Three field experiments were conducted at the Gulf Coast Research and Education Center in Bradenton, Fla., on a subsurface-irrigated EauGallie fine sand (sandy, siliceous, hyper-thermic, Alfic Hapludalf) soil.

Expt. 1. Thirty-seven ‘Mercury’ pepper transplants were set 20.3 cm apart on polyethylene-mulched, raised beds 0.76 m wide. Plots consisted of single rows 7.52 m long × 1.37 m wide with 1.62-m-long alleys and were separated by a guard row or a drainage ditch of equal width. The experiment was laid out as a randomized block design with six replications. The factorial arrangement of treatments compared two herbicides—dicamba (Banvel® 4 EC; BASF, Research Triangle Park, N.C.) and 2,4-D amine (DMA® 4; Dow AgroSciences, Indianapolis), each applied at 0, 0.11, 1.12, 11.2, and 112 g·ha⁻¹ at bloom when crown buds had just opened, or postbloom when crown fruit had attained a diameter of 2.5 to 4.4 cm. Plant vigor and leaf epinasty were rated from low to high on a scale from 0 to 10; a rating of 0 indicated dead plants or no epinasty. Fruits were harvested from 25 plants per plot on 29 Oct., and 17 Nov., then separated into marketable and cull categories and weighed.

Expt. 2. The effects of single and double exposures on ‘Mercury’ pepper plants of sublethal levels of 2,4-D or dicamba were evaluated in Spring 1987. Twenty pepper plants were transplanted 30.5 cm apart on 26 Feb. Plots were 6.1 × 1.37 m and were separated by 5.49-m alleys along the beds with two guard rows between plots. Two plants at the end of each plot were used as a guard row, so that data were collected for 16 of the 20 plants.Dicamba and 2,4-D were applied at rates of 0.01, 0.11, 1.12, 11.2 g·ha⁻¹ on 2 Apr., when flower buds were swollen, or as two sequential applications (each at full rate) on 2 and 13 Apr. (first fruits 1.3 to 1.9 cm in diameter). The experimental design was a randomized complete block with five replications. Fruits were harvested on 15 May, 3 June, and 17 June, graded, and weighed.

Expt. 3. Twenty-eight Early Cal Wonder’ pepper plants were transplanted 20.3 cm apart in plots that were 5.69 m long × 1.37 m wide on 0.76-m-wide polyethylene-mulched beds. Guard rows separated plots and there was a 4.06-m alley at the end of each plot. 2,4-D was applied at 0, 0.11, 1.12, 11.2 g·ha⁻¹ at three stages of development: prebloom (flower buds present), bloom, and postbloom (first fruits 4 cm in diameter) on 21 Mar., 29 Mar., and 15 Apr. 1988, respectively. The factorial experiment was arranged in a random-ized complete block with six replications. Yield data were collected for 20 of the 28 plants per plot (the first four plants at each end

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of the plot were not harvested), and plant height, and numbers of blooms and fruit were evaluated on one plant per plot.

Data were analyzed using the GLM procedure of SAS (SAS Institute, 1988). Orthogonal polynomials were employed to examine the nature of the response to herbicide rate. Some of the yield data were logarithmically transformed to correct for heterogeneity of variance (Snedecor and Cochran, 1980).

**Results and Discussion**

*Expt. 1.* For plant vigor, interaction between herbicide and rate was significant \(P \leq 0.001\). Higher order interactions between stage of development at application and time of evaluation after treatment were not significant; therefore, herbicide \(\times\) rate means were pooled over developmental stage and time of evaluation. Plant vigor declined more rapidly with increasing rates of dicamba than with increasing rates of 2,4-D (Fig. 1A). Interaction between stage of development at application and rate was also significant \(P \leq 0.001\); however, third- and fourth-order interactions with herbicide and time of evaluation were not. Therefore, averaged over herbicide and time of evaluation, plant vigor declined faster with increasing rates of herbicide when applied at bloom than after bloom (Fig. 1B).

Interaction between herbicide and rate was significant \(P \leq 0.05\), but higher order interactions with stage of development at application and time of evaluation after treatment were not. Therefore, data were pooled over developmental stage and time of evaluation. The increase in epinasty was greater with increasing rates of dicamba than of 2,4-D (Fig. 2A), which suggests that pepper is more sensitive to dicamba. The stage of plant development at application also influenced plant vigor; however, interaction among stage, rate, and time of evaluation was significant \(P \leq 0.0001\). The change in epinasty with increasing herbicide rate applied prebloom and at bloom is illustrated in Fig. 2B for evaluations at 2 and 4 WAT. Epinasty increased in a quadratic manner as herbicide rates were increased, but the response differed with time after treatment (Fig. 2B).

Yield responses to 2,4-D and dicamba were similar, so data were combined (Fig. 3). The effect of herbicide rate on yields of marketable and cull fruit differed with time of application \(P \leq 0.01\). With bloom applications, marketable yields decreased as herbicide rates increased from 0 to 112 g·ha\(^{-1}\) (Fig. 3A). However, postbloom application of 2,4-D and dicamba had no effect on marketable yield. Cull fruit increased as herbicide rate rose as high as 11.2 g·ha\(^{-1}\), then declined with the 112 g·ha\(^{-1}\) rate (Fig. 3B). An initial increase in yield of cull fruit also occurred with postbloom herbicide applications of 0.11 and 1.12 g·ha\(^{-1}\). However, the subsequent decline in yield of cull fruit with 11.2 and 112 g·ha\(^{-1}\), 2,4-D and dicamba was not as severe as with bloom applications. For total yield, interaction between developmental stage and rate was non-significant. Data for total yield therefore were pooled across developmental stage and a linear decline in total yield was observed with increasing herbicide rate (Fig. 3C). Averaged over rate, total yield with postbloom applications was 2 t·ha\(^{-1}\) greater than with bloom applications \(P \leq 0.02\), data not shown.

Although pepper plants appeared to be more susceptible to epinasty with postbloom applications in this experiment, the decline in vigor was less than with bloom applications, while the marketable and total yields were not affected. This suggests that the significant depression of yield by sublethal rates of auxinic herbicides may be related to their effects on reproductive development. Flower abscission and reduced fruit set in pepper and tomato have been noted with exposures to such herbicides (Gilreath et al., unpublished; Hemphill and Montgomery, 1981; Orsenigo, 1964; Robbins and Taylor, 1957).

*Expt. 2.* Dicamba and 2,4-D affected plant vigor and induced epinasty in a similar manner, so data were pooled across herbicides. When the effects of single and double applications on vigor were compared over 4 weeks following the single or initial applications, there was significant interaction between rate and number of applications \(P \leq 0.01\). This indicated that vigor declined more rapidly with the double applications than with single applications (Fig. 4A). Unlike plant vigor, the effects of herbicide rate and the number of herbicide applications on epinasty differed with time of evaluation after application \(P \leq 0.0001\). In evaluations 2 and 4 weeks after the single or initial applications, foliar epinasty increased quadratically with increasing herbicide rate (Fig. 4B). Interaction between number of applications and rate was significant \(P \leq 0.0001\) for both times of evaluation, indicating that epinasty increased more rapidly with sequential applications than with single applications.

The effect of rate on marketable yield varied with herbicide \(P \leq 0.01\) and with number of applications \(P \leq 0.05\). Marketable yield was unaffected by 2,4-D rate, but declined linearly with increasing dicamba rates (Fig. 5A). Although marketable yield was unchanged by increasing herbicide rates when applied as single applications, double applications resulted in a significant linear reduction in marketable yield as herbicide rates increased.
rate, cull yield was 85,218 kg·ha⁻¹ with single
(Fig. 6). When pooled across herbicide and
crivaled linearly with increased herbicide rates
ber of applications, cull and total yields de-
duced to 62,522 and 48,412 kg·ha⁻¹, respectively, for linear (L), quadratic (Q), or cubic (C) effects. Marketable yield was log-transformed
to correct for heterogeneity of variance. Note that vertical scales differ and that scale in (A) is
logarithmic.

Fig. 3. (A) Marketable, (B) cull, and (C) total pepper yield in response to herbicide rate and stage of
development at application (Expt. 1). Data averaged over herbicide in (A) and (B) and over herbicide
and developmental stage in (C). “” Nonsignificant and significant at $P \leq 0.05$ and $P \leq 0.01$, respectively, for linear (L), quadratic (Q), or cubic (C) effects. Marketable yield was log-transformed
to correct for heterogeneity of variance. Note that vertical scales differ and that scale in (A) is
logarithmic.

Fig. 4. Influence of single or double sprays of 2,4-D and dicamba on pepper (A) plant vigor and (B) leaf epinasty
(Expt. 2). “’” Significant at $P \leq 0.05$ and $P \leq 0.01$, respectively, for quadratic (Q) effects. Vigor was rated
from 0 to 10, with 10 being highest and 0 indicating dead plants. Rate × application means for vigor were
pooled over times of evaluation. A leaf epinasty rating of 10 indicated severe epinasty and 0 indicated none.
WAT indicates weeks after single/first application.

(Fig. 5B). Averaged over herbicide and number
of applications, cull and total yields de-
clined linearly with increased herbicide rates
(Fig. 6). When pooled across herbicide and
rate, cull yield was 85,218 kg·ha⁻¹ with single
herbicide applications and decreased to 71,112
kg·ha⁻¹ with two sequential applications ($P \leq
0.01$; data not shown). Similarly, total fruit
yield was 111,734 and 89,302 kg·ha⁻¹ with single and double applications, respectively
($P \leq 0.01$; data not shown).

Expt. 3. Plant height, averaged over rate and
time of evaluation, was 28.8 cm with
2,4-D application at bloom, which was
greater than the plant height of 22.4 cm ob-
tained with prebloom application ($P \leq 0.0001$;
data not shown). The response of plant height
to increasing rate, pooled across developmen-
tal stage and time of evaluation, was signifi-
cant ($P \leq 0.05$; data not shown): mean plant
height, which was 25.8 cm in untreated plants,
increased to 27.4 cm in plants treated with 0.11
and 1.12 g·ha⁻¹ 2,4-D then declined to 21.9 cm
with a further rate increase to 11.2 g·ha⁻¹.
The response of plant vigor to increasing
rates of 2,4-D differed with stage of develop-
ment at application ($P \leq 0.0001$). Since no
interaction occurred among rate, stage, and
time of evaluation, rate × stage means were
pooled over time of evaluation. Postbloom
applications of 2,4-D had no effect on vigor
(Fig. 7A). However, vigor decline was linear
with increasing rates of 2,4-D applied prebloom
and at bloom. Foliar epinasty increased as
herbicide rates increased, but the response
differed with stage of development at applica-
tion and the time of evaluation after applica-
tion (Fig. 7B). Epinasty in response to 2,4-D
application varied significantly with rate, de-
velopmental stage, and time of evaluation ($P \leq
0.0001$). At all times of evaluation significant
interaction occurred between rate and develop-
mental stage ($P \leq 0.0001$) and representa-
tive results are illustrated for 2 WAT (Fig. 7B).
Mild epinasty was observed with 2,4-D treat-
ment. The response to increasing rates was
quadratic and differed with developmental
stage.

For both flower and fruit number per plant
interaction among rate, developmental stage,
and time of evaluation was significant ($P \leq
0.05$). Representative data are presented for
two times of evaluation (Fig. 7 C and D). At 2
weeks after application of 2,4-D interaction
between rate and stage was significant ($P \leq
0.0001$). Flowering had just begun and even
nontreated plants had fewer than five flowers
per plant (Fig. 7C). Flower number was not
affected by prebloom application, but declined
linearly from 18 (control) to one per plant with
11.2 g·ha⁻¹ 2,4-D following bloom applica-
tions. By 4 WAT the deleterious effects of
prebloom applications were more apparent
and the linear decline in flower number was
similar to that with bloom applications.
Hemphill and Montgomery (1981) found
that 2.1 g·ha⁻¹ 2,4-D increased pepper yield
because of increased branching and flower-
ing. Bennet (1989) reported early flowering of
tomato plants treated with low concentrations
of 2,4-D, with retardation of flowering and
abnormal flowers at high concentrations. This
is consistent with the biphasic nature of aux-
ins. At low concentrations, auxins can pro-
 mote growth by cell elongation and division,
whereas, at higher concentrations, growth in-
hibition and abnormalities occur, with subse-
quent chlorosis and necrosis (Grossman, 1998).

Fewer than three fruits per plant had been
set, even on nontreated plants, 2 weeks after
prebloom or bloom applications (Fig. 7D).
Whereas a slight but significant decrease in
fruit number occurred with increasing rates of
2,4-D applied prebloom, no difference in fruit
number was obtained with bloom applica-

For both developmental stages, fruit number decreased linearly with increasing rates of 2,4-D (Fig. 7D).

For marketable and total yields, interaction between rate and developmental stage was significant ($P < 0.05$). Although both were unaffected by 2,4-D when applied postbloom, with prebloom and bloom applications there were pronounced linear decreases as 2,4-D rates increased to 11.2 g·ha$^{-1}$ (Fig. 8 A and C). Comparisons between marketable and total yields with prebloom and bloom applications revealed no significant difference due to developmental stage at the time of application (data not shown). Yield of cull fruit declined linearly with increasing rates of prebloom 2,4-D applications. However, no change in yield of cull fruit was obtained with bloom applications, and cull fruits increased with postbloom applications of 2,4-D (Fig. 8B).

Chlorosis and necrosis, which result from cyanide accumulation (Grossman, 1998), occurred only with 112 g·ha$^{-1}$, but not with lower rates. This indicates that the primary negative impact of the lower rates of 2,4-D and dicamba on pepper appears to be flower abscission, leading to reduced fruit set and yield. Epinasty and reduced plant vigor had less influence on marketable yield than did flower abscission in this study with pepper as well as in a previous study, with tomato (Gilreath et al., unpublished). Since the harvested product of pepper is the fruit, the hormonal effects on flowers and thus fruit yield are critical. Apparent stimulation of flowering at low herbicide concentrations was nonsignificant and did not translate into significantly higher yields. Low rates also had less effect on plant vigor and epinasty. As rates of dicamba and 2,4-D increased with prebloom or bloom applications or with multiple applications, increased flower abscission and reduced fruit set occurred, which were detrimental to crop yield. However, when application occurred postbloom, after most of the crop was set, sublethal rates of 112 g·ha$^{-1}$ or less did not significantly affect marketable pepper yield.

**Literature Cited**


Fig. 8. Effect of sublethal rates of 2,4-D and stage of development on (A) marketable yield, (B) cull yield, and (C) total yield of pepper (Expt. 3). **Nonsignificant and significant at $P \leq 0.01$ for linear (L) effects. Note that vertical scales differ in (A), (B), and (C).