Effect of Nitrogen Source on Pac Choi (Brassica rapa L.) Chemistry and Interactions with the Diamondback Moth (Plutella xylostella L.)

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Abstract. Two greenhouse studies were conducted to examine effects of nitrogen source on primary and secondary metabolism of pac choi (Brassica rapa L. subsp. chinensis cv. Mei Qing Choi) and diamondback moth (Plutella xylostella L.) consumption, development, survival, and body weight. Applications of a liquid organic source of nitrogen (fish hydrolysate fertilizer) were compared with a conventional fertilizer to determine whether nitrogen source directly impacts pac choi chemistry (elemental composition and phenolics) and biomass and indirectly affects diamondback moth fitness parameters. There was no significant effect of fertility treatment on pac choi chemistry or biomass with the exception of percent leaf phosphorus, which was significantly higher in the conventional fertility treatment, and p-coumarin, which was significantly higher in the organic fertility treatment. Diamondback moth also affected plant chemistry. Both calcium (Ca) and magnesium (Mg) were significantly higher in plants infested with larvae compared with uninfested plants. Fertilizer affected diamondback moth fitness with percent survival and cohort development significantly reduced on pac choi associated with the organic fertilizer. However, pac choi receiving the organic treatment was similar in regard to primary nutrients and secondary compounds compared with plants that received a conventional fertilizer.

Nitrogen (N) is an essential nutrient that crops require for growth and development (Jones, 1998; Raven and Smith, 1976). The availability and form of N may vary depending on fertilizer type. Organic fertilizers, which are derived from natural sources and approved for use in certified organic production systems, typically release N more slowly because the N is organically bound and requires mineralization before plant uptake can occur (Mengel, 1992). Conversely, synthetically derived conventional fertilizers such as ammonium nitrate, ammonium sulphate, and potassium nitrate provide N in soluble ionic forms that are readily available for uptake by plant roots (Mengel, 1992). Nitrogen supplied by fertilizers is known to influence plant composition (Penuelas and Estiarte, 1997) and thus impacts plant–insect interactions (Altieri et al., 2005; Letourneau and van Bruggen, 2006). Fertilizer effects on herbivory are associated with plant nutritional quality (proteins and amino acids) and the production of plant chemical defenses (Behmer, 2009; Sibber and Slansky, 1981). A decrease in soil N may limit protein production in plants and stimulate the production of carbon (C)-based defensive compounds (Penuelas and Estiarte, 1997). Therefore, in organic production systems, soil fertility practices have been shown to enhance plant resistance against insects, possibly by increasing C-based defenses and limiting the levels of N in tissues (Brandt and Molgaard, 2001; Geisler, 1998; Phelan et al., 1995). Conversely, by increasing the availability of soil N to plants, the ratio of C to N (C:N) in plant tissues may decrease and therefore result in limited production of C-based defenses (Folgarait and Davidson, 1995), especially those that negatively affect insect herbivores (Hsu et al., 2009; Inbar et al., 2001; Le Bot et al., 2009). These effects are generally described by the C:N balance hypothesis (Bryant et al., 1983, 1987).

Previous studies have demonstrated that fertility can affect the chemistry of plants in the brassicaeae family in ways that can impact the feeding and oviposition behaviors of a serious Brassica pest, the diamondback moth (DBM) Plutella xylostella L. (Sarfraz et al., 2009, 2010, 2011). However, it is unknown whether fertilizer inputs induce changes in the production of primary metabolites or defensive phenolic compounds, thus impacting DBM fitness and survival (Hummel et al., 2002; Staley et al., 2010). Moreover, few studies have compared the effects of fertilizer type (organic-based vs. synthetically derived) on plant chemistry and herbivory (Chen et al., 2004; Oelhaf, 1978). Implementing pest management strategies that alleviate insect outbreaks while maximizing crop production may be enhanced by understanding the effects of organic-based vs. synthetically derived fertilizers on plant–insect interactions (Altieri and Nicholls, 2003; Fageria, 2005).

This study compared N sources from an approved-for-organic-use fish hydrolysate (fish waste enzymatically broken down into peptides) with a conventional fertilizer derived from inorganic salts, which was formulated to have a similar nutrient content as the fish hydrolysate except for N form. For the fish hydrolysate, the organically bound N requires mineralization before plant uptake, whereas N from the conventional fertilizer is readily available to plants (Kristinsson and Rasco, 2000). The specific objectives of this study were to 1) assess the effect of an organic and conventional fertilizer on various pac choi (Brassica rapa subsp. chinensis cv. Mei Qing Choi) chemical composition and growth parameters; and 2) determine the corresponding effects on DBM development and survival. We chose this study system because a prior study revealed that changes in pac choi phenolic content were associated with changes in the growing environment (Zhao et al., 2009). In addition, DBM is a serious insect pest worldwide (Talekar and Lin, 1998).

Materials and Methods
Plant material and insect colony. Pac choi (Brassica rapa subsp. chinensis cv. Mei Qing

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Choi) seeds (Johnny’s Seeds, Winslow, ME) were germinated in plug trays containing a soilless growing medium, MetroMix 200 (Sun Gro Horticulture, Bellevue, WA) that consisted of Canadian sphagnum peatmoss, vermiculite, perlite, a wetting agent, and trace amounts of N:P:K. Sixty-four 2-week-old seedlings were transplanted into 16-cm black plastic containers (one seedling per container) with a soilless growing medium, Nature’s All Organic Potting Soil (Sun Gro Horticulture, Bellevue, WA) consisting of Canadian sphagnum peatmoss, composted bark, compost, and pumice. A DBM colony was established at Kansas State University (Manhattan, KS) from a cohort of individuals from Benson Research (Carlisle, PA), which was originally collected in 1988 in Geneva, NY, and maintained on a wheat germ and casein-based diet. The colony was maintained on four to five potted pac choi plants in a 0.60 × 0.60 × 0.91-m frame cage covered with 1.4-mm mesh screening in a greenhouse under natural daylight conditions and a temperature range of 20 to 23 °C. The DBM were reared on pac choi for greater than 20 generations before experiments were initiated.

**Experimental design and environmental conditions.** This study included two greenhouse experiments (spring and fall) conducted from 1 May through 5 June 2010 and 29 Oct. through 6 Dec. 2010. The experiments were conducted using a factorial treatment design with two fertility factors (conventional and organic), two DBM factors (infested and not infested), and sampling time (pre and post) for leachate analysis, in which half of the plants were destructively sampled after one application of fertilizer (pre) and the other half were sampled after three applications of fertilizer (post). Individual potted plants with treatment combinations of fertility, herbivore, and sampling time treatments were arranged in eight blocks as a randomized complete block design for a total of 64 plants. Pots were arranged on two greenhouse benches (3.6 × 9.1 m) where plants received natural daylight conditions and a temperature range of 18 to 21 °C.

To maintain uniform growing medium moisture across the blocks, an additional pac choi pot was placed at the corner of each bench, which served as an indicator container to determine when watering was needed. Pac choi plants received 750 mL of deionized water when the gravimetric weight of the indicator pots decreased by 30% from container capacity (Altimimi, 2010; Gardner, 1965) in adjacent blocks. Temperature and relative humidity were recorded using a HOBO data logger (Onset Computer Corp., Pocasset, MA) set at 30-min intervals.

**Fertility treatment.** The eight plants within each block were randomly assigned to receive either an organic or conventional fertilizer treatment (four organic and four conventional). The organic fertilizer was a soluble fish hydrolysate (Neptune’s Harvest fish hydrolysate, Neptune’s Harvest, Gloucester, MA) labeled as a 2.2N–4.3P–0.3K fertilizer and diluted to obtain a target concentration of 167 ppm N. This N concentration was selected based on previous greenhouse experiments associated with optimal N rates and pac choi yield (Altimimi, 2010). Three 100-mL diluted hydrolysate samples were submitted for nutrient testing [total N, total phosphorus (P) by potassium (K) persulfate digest, NO3-N, and NH4-N] in filtered samples by colorimetric analysis and P, K, Mg, sulfur (S), iron (Fe), sodium, and chloride (Cl) in filtered samples analyzed by inductively coupled plasma spectroscopy] by the Soil Testing Laboratory at Kansas State University (Manhattan, KS). The final concentration of the fish hydrolysate composition, based on test results, is shown in Table 1. For fish hydrolysate samples, total N was 228 mg L⁻¹ with 13% NH4-N and 0.2% NO3-N. Based on these results, the conventional soluble fertilizer was formulated using inorganic salts similar to the fish hydrolysate, except for the N composition, which is presented in Table 1. The N composition in the conventional fertilizer was 207 mg L⁻¹ total N with 29% NH4-N and 71% NO3-N. Three applications of either the conventional or organic treatment were applied during each experiment, supplying a total of 210 to 230 mg of N per container. There were three fertilizer application times: at transplant, 7 d post-transplant, and 14 d post-transplant with 330 mL of fertilizer solution applied each time.

**Leachate analysis.** Seven days after the first fertilizer application, two organically and two conventionally treated plants from each block were randomly chosen for presampling leachate and plant chemical analysis. Approximately 10 d after the third fertilizer application, final sampling of leachate, plant chemistry, and shoot biomass were evaluated. Initial and final leachate samples were processed to assess electrical conductivity (EC), pH, and macronutrient levels in the growing medium. EC is a measure of nutrients or dissolved salts in the leachate solution, recorded as dS·m⁻¹, in which a high EC value indicates a higher concentration of nutrients (Cavins et al., 2008). Containers were sampled using a pour-through method (Cavins et al., 2008), in which containers were irrigated with 750 mL of deionized water and allowed to equilibrate for 1 h over a 30-cm diameter clear plastic saucer. After equilibration, another 500 mL of deionized water was added to the containers to collect 60 mL of leachate from the saucers. Leachate samples were stored in 100-mL polyethylene vials (Fisher Scientific LLC, Denver, CO). EC and pH were measured for each leachate sample (32 initial samples and 32 final samples) using a handheld meter (Hanna Instruments, Model HI98129, Ann Arbor, MI). Leachate samples were submitted to the Kansas State University Soil Testing Laboratory (Manhattan, KS) for analysis of total N and P, and soluble K concentrations (ppm) in each sample. After digestion with K persulfate reagent in an autoclave, samples were analyzed using a Technicon AutoAnalyzer II for P content. An Alpkmek Rapid Flow Analyzer (Alpkmek Corporation, Clackamas, OR) was used to quantify NO3-N in the digested samples using the cadmium reduction method. Potassium content was determined by filtering the samples through #642 filter paper before analysis using a Flame Atomic Absorption Spectrophotometer 3110 (Perkin Elmer Corporation, Norwalk, CT).

**Herbivore treatment.** After the presampling for leachate analysis was conducted, four plants in each block, two per fertilizer treatment, were infested with 20, second instar DBM larvae 7 d after transplanting. Infestation involved the individual transfer of larvae from colony plants to the same leaf in the middle whorl of experimental plants using a fine-tipped paintbrush. To confine larvae to plants, plants were caged using 0.6 × 0.6 m nylon (2 mm mesh), which was fastened around the base of each infested pac choi plant with wooden clothespins.

At the initiation of adult emergence, moths were collected daily using the “bellows method” (Johnson, 2011). Adults were then sexed and individually weighed to the nearest 0.001 mg using an electronic balance. Male DBM were identified by the distinctive diamond-shaped patterning on the forewings, which is less evident on females (Muhammad et al., 1994; Shirai, 1993). During the emergence period, some female moths mated and laid eggs. Therefore, experiments were ended before eggs hatched and first instars began feeding, which could have affected the results. At the time of the final plant sample, counts of each larval instar as well as pupae were recorded.

**Plant chemical analysis.** Shoot biomass of aboveground plant parts were measured as fresh weight to the nearest 0.001 mg for all plants. Plant height and leaf length (cm) of one leaf in the middle whorl were recorded along with the number of leaves for all plants.

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**Table 1.** Final concentration of nutrients applied to each Brassica rapa from three applications of an organic fertilizer, Neptune’s Harvest fish hydrolysate (Neptune’s Harvest, Gloucester, MA), or three applications of a conventional fertilizer made from an inorganic salt solution computed based on chemical formulas used to compose the solution.

| Nutrient (ppm) | Organic | Conventional |
|----------------|---------|--------------|
| Total N        | 227.7 ± 14.8 | 207.4       |
| NO3-N          | 0.5     | 146.9        |
| NH4-N          | 30.3    | 60.5         |
| Total P        | 232.3   | 209.9        |
| Total K        | 60.5    | 60.6         |
| Ca             | 101.7   | 179          |
| Mg             | 11.6    | 11.7         |
| SO4-S          | 56.7    | 46.1         |
| Fe             | 0.1     | 0.3          |
| Na             | 77.7    | 77.7         |
| CI             | 32.3    | 32.6         |

*Salts used in the conventional fertilizer included: KNO3, Ca(NO3)2, MgSO4, FeETDA, NaCl, (NH4)2HPO4, and NaH2PO4.*

*Total N is shown as mean ± se to demonstrate the variability of N from three test samples of diluted fish hydrolysate.*

\[ N = \text{nitrogen}; \ P = \text{phosphorus}; \ K = \text{potassium}; \ Ca = \text{calcium}; \ Mg = \text{magnesium}; \ Fe = \text{iron}; \ Na = \text{sodium}; \ CI = \text{chlorine}. \]
For phenolic content analysis, two leaves were excised from the middle whorl (youngest, fully expanded leaves) of pac choi plants. To avoid bias from diel cycling of plant nutrients and phenolics, samples were processed between 6:00 and 8:00 AM for both experiments (You and Yang, 2001). Although greenhouse light intensities varied during the 6:00 and 8:00 AM sampling time between spring and fall experiments, the difference in light intensity was not expected to impact the comparison of plant responses between experiments (C.B. Rajeshaker, personal communication). To avoid a chemical response in plant tissue resulting from wound- ing by excising leaf tissue, leaf material was immediately frozen in liquid nitrogen and stored at –20 °C for ≈1 week until used in the analyses. For moisture content and C:N ratio, two additional leaves from the middle whorl of each plant were excised, weighed, placed in #2 brown paper bags, and stored at –20 °C until analyzed. For C and N levels, leaf material was dried in a forced-air oven set at 68 °C for 72 h and then ground in a stainless steel Wiley mill to pass through a 20 mesh screen (Scientific Apparatus, Philadelphia, PA).

Total percent C and N (both free and structural forms) were assessed from the ground tissue using a dry combustion procedure conducted by the Kansas State Soil Testing Laboratory (Manhattan, KS) through a TruSpec CN analyzer (LECO Corporation, St. Joseph, MO). Concentrations of total percent leaf P, K, Ca, Mg, S, Fe, copper (Cu), manganese (Mn), and zinc (Zn) from a 0.5-g sample of ground tissue were analyzed by an inductively coupled plasma spectrometer (SPECTRO Analytical Instruments, Kleve, Germany) after nitric acid digestion. Moisture content of the leaf tissue was assessed based on the difference between wet and dry weights of the leaf samples. Total phenolic content was analyzed using the modified Folin-Ciocalteu method (Oh, 2008; Oh and Rajashekar, 2009). Modifications included the use of 1 g of frozen leaf tissue, ground, and combined with 25 mL of 70% methanol. This extract solution was placed for 1 min in a water bath (80 °C), agitated on a shaker plate for 1 h, and filtered using No. 1 paper (Whatman, Kent, U.K.). The filtered extract (9 mL) was then evaporated to dryness by speed vacuum (Savant SVC-100H Speed Vac Concentrator; Midland, MI) under reduced pressure at 43 °C, then filtered through a 0.45-μm ascorbic acid filter (Milllex; Millipore Corporation, Bedford, MA), and re-suspended in 5 mL of 70% methanol. A 5-μL aliquot of the concentrated extract was then diluted in methanol and used in high-performance liquid chromatography for identification of the designated phenolics: chlorogenic, p-coumaric, caffeic, sinapic and ferulic acids, quercetin-3-O-glucoside, and luteolin-7-O-glucoside. Peaks for each phenolic compound were quantified and identified at 330 nm by comparing them with standard compounds of chlorogenic, p-coumaric, caffeic, sinapic and ferulic acids, and quercetin-3-O-glucoside (Sigma-Aldrich, St. Louis, MO) and luteolin-7-O-glucoside (Indofine Chemical Company, Inc., Hillsborough, NJ). These specific compounds were selected based on prior experiments with pac choi and phenolic content (M.-M. Oh, personal communication). High-performance liquid chromatography was performed by the Ruminant Nutrition Laboratory, Kansas State University (Manhattan, KS). Of the phenolics assayed, only chlorogenic, sinapic, and p-coumaric acids were detected in the leaf samples. Specific phenolic data are reported in mg/100 mL of methanolic extract.

Herbivory and herbivore response. The amount of herbivory was measured based on percent leaf area removed by feeding on plants infested with DBM larvae. For the herbivory measurement, digital images were taken of two leaves from the middle whorl (youngest, fully expanded leaves) using a camera (PowerShot SD1000; Canon, Tokyo, Japan). A photo-imaging analysis program, APS Assess Version 2.0 (APS Press, St. Paul, MN), was used to quantify total leaf area and total leaf area removed (Dudt and Shure, 1994; Lamari, 2002). Because samples contained a variable number of mixed instars, larval feeding equivalents were also used to compare herbivory between treatments. Larval feeding equivalents were calculated by dividing the percentage total leaf area consumed by the cumulative number of relative feeding equivalents. Feeding equivalents were obtained by multiplying the number of each larval instar counted on plants by the estimated relative proportion of leaf tissue required for each larval instar to complete development. Because the relative instar consumption values are unknown for DBM, we estimated consumption from those reported by Pratissoli et al. (2002) for Erinnys ello (Lepidoptera: Sphingidae). Both species are defoliating caterpillars that increase their amount of consumption incrementally from the first to fourth instars (Bellotti et al., 1992; Talekar and Lin, 1998). This procedure resulted in the following formula:

\[
PC = \frac{\sum (IN \times CR)}{CR}
\]

where PC = percentage consumption per insect per plant, PLFR = percentage total leaf area removed, IN = number of each instar, and CR = standardized consumption values for each instar (third instar = 3, fourth instar = 8, and pupa = 16).

Because there was a mixture of DBM life stages (age classes) present at the end of the experiment, mean developmental rates could not be conveniently calculated. Instead, estimates of cohort development were computed and compared between fertility treatments using degree-days (DD) as a standardized unit of comparison. Specifically, the number of individuals in each instar per plant and per treatment at the end of the experiment was multiplied by the number of DD required for that life stage to complete development based on data from Ansari et al. (2010) for DBM feeding on Brassica rapa. The total number of DD was then summed and the average number DD for the cohort was computed as follows:

\[
CDD = \frac{\sum (IN \times DD)}{TI}
\]

where CDD = average cohort DD, IN = number of each instar, DD = the number of DD needed for that life stage to complete development (third instar = 221.75, fourth instar = 159.25, pupa = 83.44, and adult = 0), and TI = total number of instars present.

Second instars were not present on plants at the end of each experiment; therefore, they were not included in the analysis. The proportion of DBM in each life stage did not differ significantly between fertility treatments within an experiment. Therefore, percent survival could be compared directly within the experiments. Percent cohort survival was computed as the sum of emerged adults and remaining larvae on plants at the conclusion of the experiment divided by the initial number of larvae (20 per plant).

Statistical analysis. Leachate N, P, K, and EC values were log10-transformed to normalize the data before analyses. Percentage data for plants (leaf tissue N, P, K, Ca, Mg, S, and moisture content) and DBM (consumption as larval feeding equivalents and cohort survival) were arcsine-transformed before analysis to normalize the data. All data presented are non-transformed.

Data pertaining to plant, insect, and leachate responses were subjected to a mixed model analysis of variance (ANOVA) using the PROC MIXED procedure (SAS Institute, 2002) with experiment (spring and fall), fertility (organic and conventional), and herbivory (plants with and without DBM) as the main effects and block as the random effect. Tests for significance were conducted for all main effects and for the two-way interactions of experiment × fertility and fertility × herbivore for plant and insect variables; and the effects of sampling time, experiment, fertility, experiment × fertility, and experiment × fertility × sampling time were used to test leachate variables. For leachate variables having significant experiment × sampling time interactions, means were then sliced to show significance for each main effect in a partitioned analysis of the least square means for the interaction (SAS Institute, 2002). The LS MEANS statement
Table 2. Analysis of variance (ANOVA) values for *Brassica rapa* response variables for block (n = 8), experiment (spring and fall), fertility treatment (conventional and organic), and herbivory (with and without *Plutella xylostella*).

| Plant variables | Block | Expt. | Fertility | Herbivory |
|----------------|-------|-------|-----------|-----------|
|                | N* df | F     | P         | N* df | F     | P         | N* df | F     | P         |
| Leaf % N       | 63  7 | 1.06  | 0.4       | 63  1 | 23.05 | <0.0001  | 63  1 | 0.4 | 0.53 | 0.01 | 0.92 |
| Leaf % C       | 62  7 | 1.16  | 0.46      | 62  1 | 113.69 | <0.001  | 62  1 | 0.89 | 0.34 | 0.21 | 0.64 |
| Leaf % P       | 62  7 | 0.57  | 0.77      | 62  1 | 55.49 | <0.001  | 62  1 | 4.29 | 0.04 | 2.46 | 0.12 |
| Leaf % K       | 62  7 | 0.88  | 0.53      | 62  1 | 47.13 | <0.001  | 62  1 | 0.32 | 0.57 | 1.43 | 0.23 |
| Leaf % Ca      | 62  7 | 0.69  | 0.68      | 62  1 | 22.1  | <0.001  | 62  1 | 0.16 | 0.69 | 5.72 | **0.02** |
| Leaf % Mg      | 62  7 | 0.18  | 0.98      | 62  1 | 4.27  | **0.04** | 62  1 | 0.07 | 0.79 | 1.72 | 0.25 |
| Leaf % S       | 62  7 | 1.49  | 0.19      | 62  1 | 9.52  | **0.003** | 62  1 | 0.28 | 0.6  | 1.27 | 0.26 |
| Leaf Cu (ppm)  | 62  7 | 4.78  | **<0.001** | 63  1 | 53.32 | **<0.001** | 62  1 | 0.11 | 0.73 | 0.02 | 0.9  |
| Leaf Fe (ppm)  | 62  7 | 1.94  | 0.08      | 62  1 | 7.26  | **0.009** | 62  1 | 1.67 | 0.2 | 0.44 | 0.51 |
| Leaf Mn (ppm)  | 62  7 | 2.24  | **0.04**  | 62  1 | 23.94 | **<0.001** | 62  1 | 0.54 | 0.22 | 1.7 | 0.19 |
| Leaf Zn (ppm)  | 62  7 | 1.31  | 0.26      | 62  1 | 88.79 | **<0.001** | 62  1 | 2.33 | 0.13 | 0.55 | 0.46 |
| Plant height (cm) | 63  7 | 0.001 | 1        | 63  1 | 4.32  | **<0.001** | 63  1 | 1.84 | 0.23 | 1.79 | 0.18 |
| Leaf number    | 63  7 | 0.001 | 1        | 63  1 | 66.11 | **<0.001** | 63  1 | 0.7 | 0.75 | 12.07 | **0.001** |
| Leaf length (cm) | 63  7 | 0.001 | 1        | 63  1 | 0.07  | 0.78  | 63  1 | 0.38 | 0.63 | 5.81 | **0.01** |
| Shoot biomass (g) | 63  7 | 0.06  | 0.99      | 63  1 | 48.3  | **<0.001** | 63  1 | 0.001 | 0.98 | 0.36 | 7.13 | **0.01** |
| Percent moisture content | 62  7 | 2.13  | **0.05**  | 62  1 | 1.86  | 0.17  | 62  1 | 0.12 | 0.73 | 0.26 | 0.61 |
| TP content b | 61  7 | 0.43  | 0.87      | 61  1 | 39.59 | **<0.001** | 61  1 | 0.12 | 0.72 | 0.21 | 0.64 |
| Chlorogenic acid b | 62  7 | 0.001 | 1        | 62  1 | 22.34 | **<0.001** | 62  1 | 0.28 | 0.6 | 0.29 | 0.59 |
| p-coumarin b | 25  7 | 4.11  | **0.01**  | 25  1 | 73.86 | **<0.001** | 25  1 | 25.56 | **<0.001** | 0.25 | 1.71 | 0.21 |
| Sinapic acid b | 54  7 | 0.66  | 0.74      | 54  1 | 4.31  | **0.04**  | 54  1 | 0.52 | 0.47 | 1.08 | 0.3  |

*bBolded values are significant at P ≤ 0.05.*

Table 3. Mean (± se) differences in *Brassica rapa* response variables between the spring and fall experiments across plants with and without *Plutella xylostella.*

| Plant variables | Plant variables | Spring | Fall |
|----------------|----------------|-------|------|
| Total % N      | 3.32 ± 0.18 b  | 4.54 ± 0.18 a |
| Total % C      | 39.06 ± 0.15 b | 41.56 ± 0.17 a |
| Leaf % P       | 0.52 ± 0.03 b  | 0.72 ± 0.02 a |
| Leaf % K       | 1.37 ± 0.10 b  | 2.80 ± 0.16 a |
| Leaf S (ppm)   | 0.56 ± 0.04 b  | 1.27 ± 0.05 a |
| Leaf Cu (ppm)  | 0.73 ± 0.14 b  | 3.52 ± 0.40 a |
| Leaf Fe (ppm)  | 166.9 ± 10.14 a| 88.43 ± 3.12 b |
| Leaf Mn (ppm)  | 114.96 ± 6.5 b | 171.51 ± 10.84 a|
| Leaf Zn (ppm)  | 41.80 ± 2.33 b | 74.74 ± 2.57 a |
| Percent moisture content | 96.44 ± 0.14 a | 96.23 ± 0.10 b |

Table 4. Mean (± se) differences for *Brassica rapa* variables between experiments (spring and fall) and herbivory (with and without *Plutella xylostella*).

| Plant variables | Spring | Fall |
|----------------|-------|------|
| Leaf % Ca      | 0.46 ± 0.40 b  | 0.66 ± 0.28 a |
| Leaf % Mg      | 0.54 ± 0.29 b  | 0.67 ± 0.16 a |
| Plant height (cm) | 14.68 ± 1.30 a| 14.43 ± 1.67 a|
| Leaf number    | 19.50 ± 3.77 a | 16.87 ± 3.26 b |
| Shoot biomass (g) | 141.93 ± 39.86 a| 119.93 ± 33.68 b|

Development, larval consumption, and survival data for either fertility treatment in each block (n = 8) were subjected to ANOVA using a PROC MIXED model for the effects of fertility, experiment, and the fertility × experiment interaction. The LS MEANS statement (SAS Institute, 2002) and Fisher’s protected LSD were used to make pairwise treatment comparisons.

To determine if experiment and fertility were correlated with various plant and insect responses, multiple regression analyses were performed using a best-subsets analysis (Minitab Inc., State College, PA) (Dallal, 2007) and the PROC REG procedure (SAS Institute, 2002). Multiple regressions applied a stepwise, backward elimination procedure (SAS Institute, 2002). Multiple regression results were assessed for best fit (linear or quadratic) in MINITAB using fitted line plots.

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Results

Plant response variables. There were no significant differences in the plant response variables between the fertility treatments in either experiment (Table 2) with the exception of percent leaf P and the concentration of p-coumarin (Table 2). In both experiments, the percentage of leaf P was significantly higher for the conventional fertility treatment (0.713 ± 0.02%) compared with the organic (SAS Institute, 2002) and Fisher’s protected least significant difference (LSD) were used to make pairwise treatment comparisons. Greenhouse environmental variables associated with light intensity, temperature, and relative humidity were also analyzed between experiments using an unpaired Student’s t test.
The concentration of p-coumarin was significantly higher in the organic fertility treatment (12.93 ± 1.18 mg/100 mL) compared with the conventional treatment (6.25 ± 0.58 mg/100 mL). Detectable concentrations of p-coumarin were found in only 19 of 32 (59%) samples in the spring experiment and six of 32 (18%) samples in the fall experiment. The experiment × fertility and fertility × herbivore interactions were not significant for any plant response variables (P > 0.05), indicating that the differences in plant responses between the fertility treatments were similar for the two experiments and for plants with and without DBM.

In general, percent leaf N, C, P, K, Ca, Mg, S, and ppm Cu, Mn, and Zn were higher in the fall experiment than the spring experiment, whereas ppm Fe, plant height, shoot biomass, and leaf number were all generally lower in the fall experiment (Tables 3 and 4). Concentrations of specific phenolics were significantly higher in the fall experiment compared with spring (P ≤ 0.05), whereas total phenolic content was significantly higher in the spring than fall experiment (P ≤ 0.05) (Fig. 1).

**Herbivory.** There was a significant experiment × herbivore interaction associated with percent leaf Ca (F = 5.53; df = 1, 59; P = 0.02). When leaf Ca was sliced for the effect of experiment, differences in Ca between spring and fall were significant (P ≤ 0.05). Slicing for the effect of herbivory (presence or absence of DBM), leaf Ca was higher in plants with DBM than in plants without DBM (P ≤ 0.0001) (Table 4). Herbivory also significantly affected percent leaf Mg (F = 4.46; df = 1, 59; P = 0.03) and shoot biomass (F = 6.21; df = 1, 53; P = 0.01), where leaf Mg was significantly higher in plants with DBM than in plants without DBM, and shoot biomass was significantly greater in plants without DBM compared with plants with DBM (Table 4).

**Diamondback moth variables.** Diamondback moth development was affected by the fertility treatments (F = 4.6; df = 1, 32; P = 0.04). Cohort development was significantly faster on plants receiving the conventional fertility treatment (83.9 ± 8.9 DD) compared with plants receiving the organic fertility treatment (105.3 ± 6.6 DD). Diamondback moth development was also affected by experiment (F = 17.13; df = 1, 32; P = 0.005) with development significantly faster in the fall (73.55 ± 9.02 DD) than the spring experiment (112.0 ± 20.46 DD). The experiment × fertility interaction was not significant (P > 0.05). Leaf consumption, based on larval feeding equivalents, was not significantly different for experiment, fertility treatment, or the experiment × fertility interaction (P > 0.05).

Percent DBM survival was significantly affected by fertility treatment (F = 5.4; df = 1, 32; P = 0.02). Specifically, for both the spring and fall experiments, percent survival was significantly higher on plants receiving the conventional fertility treatment (64% ± 0.04%) compared with plants receiving the organic fertility treatment (46% ± 0.05%). Percent survival was also significantly affected by experiment (F = 5.37; df = 1, 32; P = 0.02) although the experiment × fertility interaction was not significant (F = 1.35, df = 1, 16; P = 0.25), which suggests that the effects of fertility and time the experiment was conducted on survival were independent. Regression analysis revealed that DBM survival varied significantly with leaf Mg, in which survival was negatively related to leaf Mg at concentrations greater than 0.6% in the fall experiment (F = 6.79; df = 1, 16; P = 0.02). Changes in leaf Mg may be responsible for the high percentage variation in survival for the fall experiment (R² = 97%) (Fig. 2). However, the regression of survival and leaf Mg was not significant in the spring experiment (F = 4.07; df = 1, 16; P = 0.06).

Fertility treatment did not significantly affect male or female DBM body weights nor was there a significant experiment × fertility interaction (P > 0.05). Female body weights were significantly affected by experiment (F = 12.43; df = 1, 16; P = 0.002), in which body weights were higher in the spring experiment (4.87 ± 1.10 mg) than the fall experiment (2.73 ± 1.50 mg). Male body weights were not significantly affected by experiment (P > 0.05).

**Leachate variables.** There were no significant differences in leachate N, P, K, pH, and EC between the fertility treatments (Table 5). However, leachate N, P, K, pH, and EC were significantly affected by sampling time (Table 5), in which all variables were significantly higher at pre-sample (before introducing DBM) compared with the post-sample (end of experiment). For leachate P, the pre- and post-sample concentrations (mean ± SE) were 30.8 ± 1.7 ppm and 18 ± 2.5 ppm, respectively. For the other leachate variables (N, K, pH, and EC), there was a significant experiment × sampling time interaction (Table 5), indicating that the magnitude of difference between the pre- and post-samples was not the same for the spring and fall experiments. Slicing tests for main effects (experiment and sampling time) were used to determine the significance of each effect while holding the other constant. Slicing for the effect of experiment across sampling times revealed that leachate N, K, and pH were all significantly different between the spring and fall experiments (P < 0.0001), whereas slicing for sampling times across experiments revealed that leachate N and K were not significantly different between sampling times (P > 0.05) but leachate pH was (P < 0.0001). For leachate EC, slicing for sampling times across experiments showed that the EC was significantly different across sampling times between experiments (P < 0.0001), and the EC was significantly different between sampling times when experiment was held constant (P < 0.001). Using pairwise treatment comparisons, leachate N
and K were significantly greater at the pre-sampling time in the spring experiment, leachate pH was significantly higher in the spring post-sampling time, and leachate EC was significantly higher at the spring pre-sampling time compared with other sampling time and experiment combinations (Fig. 3).

Greenhouse environmental conditions (light intensity, temperature, and relative humidity) were significantly higher in the spring experiment than in the fall experiment. Greenhouse conditions were significantly different between experiments, including light intensity ($t = –8.69$, $df = 856$, $P \leq 0.001$), which was higher in the spring experiment ($18,324 \pm 376$ lumens/ft$^2$) compared with the fall experiment ($12,825 \pm 505$ lumens/ft$^2$). Temperature was also significantly ($t = –18.62$, $df = 2613$, $P \leq 0.001$) higher in the spring experiment ($25 \pm 0.1 ^\circ C$) compared with the fall experiment ($21 \pm 0.1 ^\circ C$); and percent relative humidity was significantly higher ($t = –39.78$, $df = 2613$, $P \leq 0.001$) in the spring experiment ($59\% \pm 0.4\%$) than the fall experiment ($33\% \pm 0.3\%$).

**Discussion**

In this study, the type of fertilizer used had no effect on any of the leachate variables and only a minimal effect on pac choi chemistry. With respect to plant responses, percent leaf P was significantly higher in the conventional fertility treatment than the organic fertility treatment in both experiments. Other studies examining plant P content in relation to different types of fertilizers are not known for pac choi. The protein competition model proposed by Jones and Hartley (1999) suggests that P has a limiting effect on plant growth and C fixation through reduced phosphorylation. Therefore, it might be expected that the higher levels of P in conventionally fertilized pac choi may have limited available C for use in accumulating phenolics. However, there were no differences in total phenolic content between the fertility treatments and no differences in any of the plant growth responses. These results differ from Zhao et al. (2009), who examined nutrient and phenolic contents in greenhouse-grown pac choi (*Brassica rapa* subsp. chinensis ‘Mei Qing Choi’) and compared an organic fish hydrolysate fertilizer with a conventional fertilizer (inorganic salt solution formulated similar to fish hydrolysate). They found increased phenolic contents in plants that received the fish hydrolysate fertilizer compared with plants that received the conventional treatment. It must be noted, however, that Zhao et al. (2009) applied higher rates of N and added a slow-release compost amendment to the fish hydrolysate treatment, which may have immobilized and limited N provided to plants by the fish hydrolysate. Based on the C:N balance hypothesis (Bryant et al., 1983), it would be expected that leaf C and phenolic content would increase when N resources were limited for plant growth and development.

In this study, the organic fertility treatment was designed to create a limited N supply for pac choi plants relative to the conventional fertility treatment. However, based on the plant responses, it appeared that N was not actually a limiting factor in the organic fertility treatment nor were there differences in leaf C and phenolic content between the treatments. The similarities between the fertilizer treatments in this study may be attributed to rapid mineralization of organically bound N, which is common for animal-based fertilizers (Hartz and Johnstone, 2006). Moreover, some greenhouse studies have reported similar plant chemistries and growth responses in plants receiving fish-based and synthetically based fertilizers (Aung and Flick, 1980; Emin, 1981).

Diamondback moth herbivory impacted plant chemistry with larval feeding, possibly increasing the levels of leaf Ca and Mg in both experiments. Because Ca and Mg are n

Table 5. Analysis of variance (ANOVA) values for block ($n = 8$), experiment (spring and fall), sample time (ST) (pre-sample: before *Plutella xylostella* introduction, and post-sample: completion of experiment), and the experiment $\times$ ST interaction associated with the leachate variables.$^a$

| Main effects$^a$ | N (ppm) | F | df | P | P (ppm) | F | df | P | K (ppm) | F | df | P | pH | F | df | P | EC$^a$ | F | df | P |
|------------------|---------|---|----|---|---------|---|----|---|---------|---|----|---|-----|---|----|---|------|---|----|---|
| Block            | 0.76    | 7 | 0.62 | 0.65 | 0.71 | 1.1 | 1 | 0.37 | 1.99 | 1 | 0.06 | 3.57 | 1 | 0.001 |
| Experiment       | 0.01    | 1 | 0.93 | 0.001 | 1 | 0.98 | 0.01 | 1 | 0.92 | 4.71 | 1 | 0.03 | 95.29 | 1 | <0.001 |
| ST               | 65.56   | 1 | <0.001 | 57.47 | 1 | <0.001 | 94.42 | 1 | <0.001 | 138.2 | 1 | <0.001 | 48.54 | 1 | <0.001 |
| Experiment $\times$ ST | 6.45 | 1 | <0.001 | 3.58 | 1 | 0.06 | 6.59 | 1 | 0.01 | 11.95 | 1 | <0.001 | 22.41 | 1 | <0.001 |

$^a$There were 126 plants used with four containers per fertility treatment in each of eight blocks at two sampling times for the two experiments.

B Bolded values are significant at $P \leq 0.05$.

$^x$Reported in mhos/cm.

N = nitrogen; P = phosphorus; K = potassium; EC = electrical conductivity.
associated with plant defenses related to stress, it may be expected that levels of both nutrients would increase in response to DBM herbivory (Kolupaev et al., 2008; Rodriguez-Serrano et al., 2009). Likewise, Sarfraz et al. (2009) found significantly higher percentages of Ca and Mg in the leaf tissues of *Brassica napus* that had been infested with DBM compared with uninfested plants. With respect to insect defense, Sarfraz et al. (2009) also found that plants with higher levels of Ca and Mg resulted in longer development times and shorter adult lifespans of DBM.

In both experiments, DBM had a significantly higher survival rate and developed faster on plants treated with the conventional fertilizer. Although neither larval consumption nor adult body weight were affected by the type of fertilizer treatment, our findings suggest that plant factors associated with the organic fertilizer treatment were less suitable for DBM compared with the conventional fertilizer treatment. However, there is no clear explanation for the observed differences in development and survival. The impact of different greenhouse conditions (light, temperature, and relative humidity) is not known on DBM fitness in this study but significantly varied between experiments. It is possible that the lower level of P in pac choi leaves in the organic fertilizer treatment adversely affected DBM. For example, P availability during larval development may be an important factor influencing insect life history parameters (Visanuvimol and Bertram, 2011), and decreased dietary P has been shown to reduce the growth rate of *Choristoneura occidentalis* and *Manduca sexta* larvae (Clancy and King, 1993; Perkins et al., 2004). However, DBM development was not correlated with leaf P in the current study. Alternatively, treatment differences in other unmeasured plant components such as amino acid composition and concentration or changes in glucosinolate content may have affected the nutritional quality of pac choi for DBM. For example, Bejai et al. (2010) found specific glucosinolate levels (1-methoxyindol-3-methylglucosinolate, 3-benzoyloxypropylglucosinolate and 4-hydroxybenzylglucosinolate) were associated with increased DBM pupal weight and prolonged larval development in *Arabidopsis thaliana*. We did not assess specific glucosinolate levels in our study as a result of constraints of time and plant tissue required for analysis. Furthermore, changes in total glucosinolate content in *Brassica* plants have not been found to be associated with DBM larval performance (Arany et al., 2008; Ratzka et al., 2002).

As a result of the difficulties in managing DBM in crop production systems (Eigenbrode and Shelton, 1990), the current study contributes information that may be useful to producers. First, both fertility treatments appeared to provide comparable N sources for growth of pac choi. When implementing an organic fertilizer program, producers may choose fish-based fertilizers that deliver N to plants similar to conventional fertilizers, which is different from slower-release plant-based fertilizers and compost amendments. As such, future research, evaluating the use of multiple organic-based materials, is warranted. Second, it appears that the effect of the organic fertilizer treatment negatively impacted DBM development and survival. Although general predictions regarding pac choi susceptibility to DBM feeding under organic fertilizer programs compared with conventional fertilizer programs cannot be made at present, further studies may demonstrate that crop susceptibility changes may be associated with the type of fertilizer used. The possibility that the fish hydrolysate fertilizer induced pac choi defenses against DBM should be examined further by focusing on leaf P levels or other components affiliated with the hydrolysate formulation not measured in this study.

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