Phosphorus, Zinc, and Boron Influence Yield Components in ‘Earliglow’ Strawberry

Geoffrey M. May and Marvin P. Pritts
Department of Fruit and Vegetable Science, Cornell University Ithaca, NY 14853

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Abstract. The main effects and interactions of soil-applied P, B, and Zn on yield and its components were examined in the field at two pH levels with ‘Earliglow’ strawberries (Fragaria × ananassa Duch.). Applied nutrients had significant effects on several yield components, but responses depended on the levels of other nutrients or the soil pH. At a soil pH of 5.5, yield responded linearly to B and quadratically to P. At pH 6.5, P interacted with B and Zn. Fruit count per inflorescence was the yield component most strongly associated with yield followed by individual fruit weight. However, these two yield components responded differently to soil-applied nutrients. Foliar nutrient levels generally did not increase with the amount of applied nutrient, but often an applied nutrient had a strong effect on the level of another nutrient. Leaf nutrient levels were often correlated with fruit levels, but foliar and fruit levels at harvest were not related to reproductive performance. Our study identifies some of the problems inherent in using foliar nutrient levels to predict a yield response and demonstrates how plant responses to single nutrients depend on soil chemistry and the presence of other nutrients.

Achieving the optimum level of nutrition for horticultural crops is a major objective of practitioners. Realization of this objective is complicated because: 1) at least 12 essential nutrients are involved, 2) many environmental factors influence the availability and uptake of these nutrients, 3) genetic differences in uptake and usage exist at all taxonomic levels, and 4) nutrients interact among themselves.

The first three items have been explored extensively for most major crops, including strawberries (May and Pritts, 1990). However, investigations of nutrient interactions in the field have been lacking because of the difficulty in testing for such effects. For example, scientists often use factorial experiments to test for interactions. An experiment designed to quantify interactions among three factors at five levels would result in 125 treatment combinations if a complete factorial design was used. When replication is introduced, the number of experimental units becomes unmanageable.

Response surface methodology has been developed to enable the researcher to minimize the number of treatment combinations for the generation of a multidimensional response (Box and Draper, 1987). Using the above example, only 15 treatment combinations are needed to generate a response surface, assuming no third order or higher responses or interactions exist (Cochran and Cox, 1957). The ability to use fewer treatment combinations to generate response surfaces allows the study of nutrient interactions in field situations.

Strawberries are an ideal model for nutrient interaction studies in perennial crops. First, they are relatively precocious, producing their first crop within months of planting. Second, the plants are small, so that many plots can be established within uniform soils. Third, they can be easily excavated and divided into vegetative and reproductive components, allowing for detailed evaluation of observed growth or yield responses. Fourth, when compared to other perennial fruit crops, cropping history has little influence on flower bud initiation and fruiting.

The objective of our research was to evaluate the main effects and interactions among P, Zn, and B as they affect tissue nutrient status and yield in ‘Earliglow’ strawberries. These three nutrients were chosen for several reasons. First, a negative P-Zn interaction has been reported in other crops (Barnette et al., 1936; Murphy et al., 1981; Olsen, 1972; West, 1938). Second, a survey of leaf analysis results found that Zn and B were consistently low in strawberry plantings, and P was often excessive (Mitlehner, 1989). Our review of results from Cornell Univ.’s Plant Tissue Analysis Laboratory, Ithaca, N.Y., reflected a similar pattern. Third, those who grow strawberries in matted-rows often use a P-containing fertilizer at renovation, even though P levels are usually amended at planting. If P does interact negatively with these other nutrients, then fertilization practices at renovation may require modification.

A second objective was to determine if main effects and interactions were consistent at different soil pH levels. Soil pH is known to have a major effect on nutrient availability, and since commercial fields vary considerably in pH, it is important to know how widely applicable results are from experiments conducted under a single set of conditions.

We selected a soil for the experiment that was not deficient in P, B, or Zn, a situation typical of most soils on which strawberries are grown in the northeastern United States. However, we used a wide range of nutrient additions to generate imbalances that might occur in a commercial situation, recognizing that responses would likely be different (and possibly stronger) on nutrient-poor soils.

Materials and Methods

Interactions among P, B, and Zn at two pH levels in ‘Earliglow’ strawberry were examined at Cornell Orchards, Ithaca N.Y. The soil was a Collamer silt loam (Typic Hapludalf, pH = 5.5). Soil testing before treatment application, using an ammonium acetate extraction procedure, indicated about (kg-ha−1) 11 P, 92 K, 2800 Ca, 435 Mg, 159 Fe, 139 Mn, 2.2 Zn, 0.2 B, and 6.7 Cu.

A three-factor, rotatable, central composite design (Cochran and Cox, 1957) was used to identify the specific orthogonal treatment combinations required to obtain a second-order re-
Nutrients were soil-incorporated before planting. Rates of application ranged from 11 to 123 kg P/ha as triple super phosphate, 2.8 to 45 kg Zn/ha as zinc sulfate, and 0.2 to 4.5 kg B/ha as sodium octaborate. Ground agricultural limestone was used to adjust soil pH level in half of the plots from the existing pH of 5.5 to 6.5.

Applications of P, Zn, B, and lime were made in Oct. 1986 before rototilling of each plot. To ensure uniform distribution, zinc sulfate and sodium octaborate were first dissolved in water, before rototilling of each plot. To ensure uniform distribution, these nutrients were then applied to each plot using a hand sprayer. Other than N, no additional nutrients were applied during the course of the experiment. Subsequent soil testing (ammonium acetate extraction) was conducted in Spring and Fall 1987 and in Spring 1988.

On 23 Apr. 1987, certified nursery plants of ‘Earliglow’ were planted 30 cm on square, forming a 40-plant grid in each plot. Standard cultural practices were followed (Pritts et al., 1988), with the exception that runners were removed so plants did not form a solid bed. Napropamide at 2.2 kg·ha⁻¹ was applied 2 weeks after planting and again at 4.5 kg·ha⁻¹ before straw mulch application in November. Ammonium nitrate was applied 6 weeks after planting to supply 34 kg N/ha, and calcium nitrate was applied in early September to supply 22 kg N/ha. Soil moisture was maintained near field capacity throughout the study with overhead irrigation, as indicated by soil moisture tensiometers.

Runners were removed weekly to maintain a constant plant density within each plot. Entire plants were excavated beginning 2 June 1987 and continuing at 30-day intervals throughout the first year of establishment (1987) and first fruiting season (1988). Plants were separated into roots, crowns, petioles, leaf blades, and fruit, washed with tap water, then rinsed with distilled water. Plant material was oven-dried at 65°C, weighed, and ground in a Wiley mill to pass through square grids 1.4 × 1.7 mm.

Nitrogen concentration in leaf lamina was determined by a Kjeldahl procedure. The P, K, Ca, Mg, Fe, Zn, B, Mn, Cu, and Mo content was determined by inductively coupled plasma spectroscopy (Model 975 Plasma Atomcomp ICP Spectrograph with ICAP 61 Update, Jarrell-Ash, Pittsburgh). Leaf area was determined using a portable area meter (LI-COR Model LI-3100, Lincoln, Neb. Aboveground plant dry weight, number of branch crowns and fruit per inflorescence, inflorescence length, fruit count per plant, fruit weight, achenes per square centimeter of surface, and fruit weight per achene were recorded at each sampling date. Yield component data were pooled for each plot. A final leaf sample for tissue nutrient analysis was collected ≈5 weeks after postharvest leaf removal. Fruit soluble solids concentration (SSC) was determined with a hand-held refractometer (Thomas Scientific, Swedesboro, N.J.).

Statistical analysis of the central composite design was used to derive meaningful regression equations that accurately described the response surfaces for the variables of interest. Values for nutrient concentrations were transformed before analysis by first subtracting the mean, then dividing by it, resulting in a scale from -1 to 1 for each nutrient. Using a transformed scale in regression analysis has three advantages over using the actual levels of applied nutrient. First, regardless of the nutrient involved, -1 represents a low level and +1 a high level. This scheme is important when the differences in ranges between low and high levels are very large (as was the case with P and B). Second, the transformed variables are orthogonal, allowing unbiased estimates of regression coefficients to be calculated. Third, regression coefficients among components of the model can be compared directly with each other. Also, this standardized scale can be converted easily to the actual applied values through multiplication.

These new, orthogonal variables were used in regression analysis to calculate the response surface, using the following full model:

\[
\text{Response} = \text{Block} + P + Zn + P \times Zn + P \times B + B \times Zn + P^2 + B^2 + Zn^2.
\]

Sequential sums of squares were accumulated for the full model. Single-degree-of-freedom F tests were then used to evaluate the significance of the individual components of the model. For the generation of response surfaces, new models were constructed using only the significant terms from the full model, with the exception of fruit count per inflorescence, where the full model was used (since the full model was significant, but individual coefficients were not).

### Results

Preplant, soil-applied nutrient additions of P, Zn, and B were correlated with soil test results in each of the 2 years (Table 2). A decrease of ≈0.5 pH units was measured during the course of the experiment in plots that initially received lime applications. Plant growth variables were unaffected by liming; however, yield components were affected differently at pH 5.5 and 6.5, so results are discussed separately.

**Results at pH 6.5.** Yield increased with increasing B at a high P level but decreased at a low P level (Table 1, Fig. 1 top).

**Table 2.** Significant (P < 0.01) correlation coefficients (r) between mineral nutrient treatment levels (1986) and soil test values (1987 and 1988).

| Soil test  | P   | Zn  | B   |
|-----------|-----|-----|-----|
| Spring 1987 | 0.94 | 0.97 | 0.84 |
| Fall 1987  | 0.74 | 0.65 | NS  |
| Spring 1988 | 0.55 | 0.93 | 0.71 |

NS: Non-significant.
Fig. 1. Yield response (grams per six plants) to P and B at pH = 6.5. For scale conversion, units of P range from 11.2 (–1) to 123.3 (1) kg·ha\(^{-1}\) and B from 0.2 (–1) to 4.5 (1) kg·ha\(^{-1}\). (Top) Zn = 1.0 (23.9 kg·ha\(^{-1}\)). Yield = 1470 + 605(P*B) - 394(B*Zn), \(R^2 = 0.213, P < 0.007\). (Bottom) Zn = -1.0 (2.9 kg·ha\(^{-1}\)). Yield = 1470 + 605(P*B) + 394(B*Zn), \(R^2 = 0.213, P < 0.007\).

Zinc moderated this response (Fig 1 bottom), reflecting an interaction of P with B and Zn (Table 3). Fruit count per inflorescence was the only yield component correlated strongly with yield \(r = 0.62, P < 0.01\), although fruit size and yield were marginally correlated \((r = 0.29, P < 0.06)\). The response surface for fruit count per inflorescence resembled that of yield, but the regression coefficients were not statistically significant (Fig. 2). Branch crown count was significantly affected by P and B but exhibited a very different response surface than that for yield (Fig. 3). Nutrient treatments had no effect on fruit SSC, fruit size, achene spacing, or inflorescence count, although inflorescence length was associated positively with a wider achene spacing on the fruit receptacle \((r = 0.34, P < 0.02)\).

Foliar levels of P, Zn, and B fluctuated during the season (Fig. 4), and by renovation in 1988, only P was still influenced significantly by the soil nutrient addition made in 1986 \((r = 0.29, P <0.05)\). Leaf P generally increased with soil P >30 kg·ha\(^{-1}\), but decreased as soil B increased (Fig. 5). Leaf B and Zn, however, were positively influenced by the level of applied P \((P < 0.02)\) but not by the level of applied B or Zn.

None of the foliar nutrient levels at renovation were correlated

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Table 3. Yield components and reproductive variables with significant regression coefficients for the full response surface model.

| Variable                        | Independent                                                                 |
|---------------------------------|-----------------------------------------------------------------------------|
| pH 6.5                          | \(Yield = 606(P \times B)^{**}, -395(P \times Zn)^*\)                        |
| Branch crowns/plant             | None                                                                        |
| Inflorescences/plant            | None                                                                        |
| Fruit/inflorescence             | None                                                                        |
| Individual fruit wt             | None                                                                        |
| Achenes/fruit                   | None                                                                        |
| Achene density                  | None                                                                        |
| Soluble solids concan           | None                                                                        |
| Runners/plant                   | None                                                                        |
| Aboveground plant wt            | 12.2(B^2)*, -1.35(Zn^2)*                                                   |
| pH 5.5                          | \(Yield = 167(B)^*, 658(P^2)^{**}\)                                        |
| Branch crowns/plant             | 4.36(P \times B)^{***}, 6.09(P \times Zn)^{***}                           |
| Inflorescences/plant            | None                                                                        |
| Fruit/inflorescence             | None                                                                        |
| Individual fruit wt             | 0.368(P)^*, 0.349(B)^*                                                     |
| Achenes/fruit                   | None                                                                        |
| Achene density                  | None                                                                        |
| Soluble solids concan           | None                                                                        |
| Runners/plant                   | -31.4(Zn)^{**}, 30.1(Zn^2)^{**}                                            |
| Aboveground plant wt            | 210(P)^{**}, 200(P \times Zn)^{**}                                         |

**,**,** Probability of significance is \(P < 0.10, 0.05, \text{ or } 0.01\), respectively.
with yield. For K, P, Ca, Mn, Cu, and Zn, foliar and fruit levels were correlated positively, but no correlation existed for levels of Mg, Fe, or B. Plant size was not correlated with yield.

Results at pH 5.5. Strawberry yields were significantly affected by B (Fig. 6), but the interaction of B with P or Zn was not significant (Table 3). Boron applications increased yield regardless of P and Zn levels. As at the higher pH, fruit count per inflorescence ($r = 0.65$, $P < 0.001$) and individual fruit weight ($r = 0.31$, $P < 0.04$) were correlated with yield, and their responses to nutrient additions somewhat paralleled the yield response (Figs. 7 and 8). Again, although the response of fruit count per inflorescence was very similar to that of yield, regression coefficients that describe the response of this variable to nutrient additions were not significant (Fig. 8).

Fig. 3. Response of branch crowns per plant to P and B at Zn = 0 (23.9 kg·ha$^{-1}$) and pH = 6.5. Branch crown count = 11.0 - 3.15(P)$^2$, $R^2 = 0.09$, $P < 0.05$. For scale conversion, units of P range from 11.2 (-1) to 123.3 (1) kg·ha$^{-1}$ and B from 0.2 (-1) to 4.5 (1) kg·ha$^{-1}$.

Fig. 4. Seasonal pattern of P, B, and Zn concentration in leaf lamina of plants set in Apr. 1987 and harvested over the following 16 months for the central treatment of the design of the experiment (kg·ha$^{-1}$: 67 P, 23.9 Zn, and 2.35 B).

Fig. 5. Response of leaf P at renovation to P and B at Zn = 0 (23.9 kg·ha$^{-1}$) and pH = 6.5. Percent leaf P = 0.201 + 0.014(P) + 0.034(P)$^2$ - 0.013(B) - 0.03(B$^2$Zn), $R^2 = 0.339$, $P < 0.002$. For scale conversion, units of P range from 11.2 (-1) to 123.3 (1) kg·ha$^{-1}$ and B from 0.2 (-1) to 4.5 (1) kg·ha$^{-1}$.

Fig. 6. Yield response (grams per six plants) to P and B at Zn = 0 (23.9 kg·ha$^{-1}$) and pH = 5.5. Yield = 1427 + 410 (P$^2$) + 166(B), $R^2 = 0.153$, $P < 0.03$. For scale conversion, units of P range from 11.2 (-1) to 123.3 (1) kg·ha$^{-1}$ and B from 0.2 (-1) to 4.5 (1) kg·ha$^{-1}$.

Nutrient additions had no effect on inflorescence count, fruit SSC, or achene spacing (Table 3).Inflorescence length, however, was positively correlated with fruit size ($r = 0.39$, $P < 0.009$), yield ($r = 0.36$, $P < 0.02$), achene spacing ($r = 0.34$, $P < 0.02$), and the level of applied B ($r = 0.31$, $P < 0.04$).

Leaf P at renovation showed a quadratic response to soil-applied P (data not shown), similar to that at the higher pH (Fig. 5). Leaf B exhibited a quadratic response to P ($P < 0.06$) but not to B additions (data not shown).

Leaf Zn at renovation was negatively correlated with yield ($P < 0.06$), but correlations of yield with other foliar nutrient levels at renovation were not significant (data not shown). Fruit and leaf nutrient levels were correlated positively for P, Mn,
That we observed could have been influenced by the Ca and Mg levels of soil pH with a limestone amendment. The responses we observed rather, we will make three major points that do not depend on knowing the exact nutrient status of the soil or the response of plants to individually applied nutrients from deficient through toxic levels. These points are as follows: 1) soil pH can alter plant response to applied nutrients; 2) predictions about plant response to applied nutrients are difficult when interactions with other nutrients exist; and 3) individual yield components respond differently to applied nutrients, implying that there may be more than one optimal combination of nutrients for maximum yield.

Nutrient interactions and soil pH influenced plant response to individual nutrients. Yield and certain yield components exhibited a strong response to soil-applied B (Figs. 1, 6, and 8). However, the yield response of strawberries to soil-applied B depended on the level of soil P and pH. Yields at the higher pH (6.5) varied among treatments nearly 2-fold, due primarily to interactions of P with B and Zn. Yield at the lower pH was also influenced by P and B but not by an interaction between the two nutrients (Fig. 6).

The existence of nutrient interactions has been well documented (Adams, 1980; Kirsch, 1959; Murphy et al., 1981; Robson and Pitman, 1983). For example, detrimental effects of P fertilization on Zn-deficient soils have been reported for many crops (Olsen, 1972). Efforts to uncover the mechanism of the P-Zn interaction have led to the study of P-Zn reactions in the soil and reactions within the plant involving uptake, translocation, metabolism, and growth. Temperature and Fe levels have been found to influence P-related Zn disorders (Adams, 1980).

Phosphorus can also interact with B. Boron is essential for root-tip elongation which, in turn, affects P uptake (Pollard et al., 1977). However, P also influences B uptake. Bingham and Garber (1960) and Bingham et al. (1958) reported that excessive P additions reduced uptake of B in citrus. Boron and phosphates are also involved in complex adsorption and precipitation reactions with sesquioxides and clay minerals in the soil. Thus, the potential for interactions exists at many levels, and the expression of these interactions may depend on environmental conditions or the presence of other nutrients.

Blatt (1976, 1982), working with ‘Midway’ strawberry on an acidic sandy soil in Nova Scotia, investigated possible P-B interactions under field conditions. Using a factorial approach, he found that neither initial vegetative growth nor subsequent fruit yields were affected by P or B additions and that there was no P-B interaction. We also found no P-B interaction at a low pH.
but we did find a yield response to B that was mediated both through fruit count per inflorescence and fruit size. Although the response of fruit count per inflorescence to B was not statistically significant, this yield component was highly correlated with yield at both pH levels, and the response surfaces were similar (Figs. 1, 2, 6, and 7).

Boron influences auxin activity (Pilbeam and Kirkby, 1983), and since auxin is related to both inflorescence development and fruit receptacle expansion (Archbold and Dennis, 1985), one would expect yield to respond to supplemental B if its ambient levels were inadequate and P was adequate. Inflorescence length may be a useful indicator of auxin activity, as this variable was associated with achene spacing on the receptacle and was positively associated with yield, fruit size, and applied B at the lower pH.

Interactions among nutrients may help explain inconsistent responses to applied nutrients that have been reported in the literature. Neilson and Eaton (1983) studied the effects of B nutrition on strawberry yield components using solution culture and found no yield response to B applications from 0.01 to 1.25 ppm. Riggs et al. (1987) found that B applications up to 4.5 kg ha⁻¹ decreased yields of ‘Tristar’ and ‘Benton’ strawberries on Quatama loam soils suspected to be low in B in the Willamette Valley, Ore. In our study, at a soil pH of 5.5, yield responded positively to B. However, at a soil pH of 6.5, the responses to applications of B were either positive, neutral, or negative, depending on the specific level of P fertilization (Fig. 1).

Our results demonstrated an important shortcoming of field studies that examine single nutrients, without considering interactions with other nutrients. Adams (1980), Kirsch (1959), Murphy et al. (1981), and Sumner and Farina (1986) emphasized the importance of designing nutritional experiments that permit measurements of potential interactions to avoid possible misleading main-effect conclusions.

While it is important for horticulturists to attempt to identify the nutritional conditions that optimize total yield, it is likely that all of the various components contributing to final yield do not have the same nutritional requirements. For example, number of branch crowns, fruit count per inflorescence, fruit weight, and runner production responded differently to additions of P, Zn, and B. Also, responses changed with soil pH. These observations and the finding of interactions suggest that there may not be a single optimum combination of nutrients that maximizes yield. Yield optimization, in reality, likely consists of multiple sets or nutrient levels or ranges, which of themselves are not limiting to plant growth or yield.

There are many inconsistencies in the literature describing the response of strawberries to fertilization (May and Pritts, 1990). Some of these inconsistent responses may reflect nutrient interactions such as those found in our study, while others may reflect variations in climate, soil moisture, soil type, plant condition, and cultural practices. Tissue analysis has been viewed as an invaluable tool to assess the nutritional status of perennial crops and to directly evaluate plant response across a wide range of conditions (Bould, 1964). However, in our study, leaf nutrient levels during the recommended postharvest period in 1988 generally were not related to yield, nor were foliar nutrient concentrations in 1988 correlated with nutrient levels in the soil.

At pH 6.5, applied P was correlated with leaf P, which in turn was correlated with fruit P. Applied Zn was not correlated with foliar Zn, although foliar and fruit Zn levels were correlated. No correlation existed among soil, foliar, or fruit B or between yield and fruit P, B, or Zn. Further, B levels in ripening fruit (17 to 22 ppm) were much lower than foliar levels at the same time (54 to 73 ppm). Therefore, practices that increase soil nutrient levels may not lead to increased foliar levels, especially if nutrient dilution occurs as growth is stimulated. In addition, a strategy that successfully increases foliar nutrients may not always lead to higher levels in reproductive tissues. Finally, higher levels in reproductive tissues may not be associated with higher yield or fruit quality.

Others have concluded that foliar analyses are not particularly useful indicators of nutrient status for particular elements. Riggs et al. (1987) found that soil applications of B were not effective in increasing total B in reproductive tissues and that foliar B levels were a poor predictor of levels in reproductive tissues.

A plant nutrient analysis alone may not be sufficient to identify treatments that will increase yield economically, at least when plants are grown on a reasonably fertile soil. The challenge of understanding nutritional responses is great, considering the thousands of possible interactions that could occur among the 12 mineral nutrients for a single variable under a single set of conditions. Adding to the complexity of evaluating plant nutrient status is the observation that leaf nutrient levels fluctuate seasonally (Fig. 4). Even if one could find specific times when the level of certain nutrients is predictive of plant performance, growers may not accept the time and expense involved with multiple sampling times.

Finally, the results of our study are limited because we examined the response of only a single genotype. Some growers have reported that ‘Earliglow’ is relatively tolerant to low nutrient levels, but this evidence is anecdotal. With many crop plants, including strawberries (John et al., 1976; Peterson et al., 1986; Renquist and Hughes, 1985), significant genetic variation exists in the ability to extract or absorb nutrients from the rhizosphere or tolerate low levels of nutrients. More work will be required to identify genotypes tolerant to nutrient imbalances.

Nutrient additions significantly affected strawberry yield through various yield components, and these responses were pH dependent. Interactions among applied nutrients also occurred, making it difficult to predict strawberry plant response under different conditions. Soil type, climate, cultivar, plant condition, cultural practices, and the levels of other plant nutrients also likely influence the response of the strawberry plant to P, B, and Zn. Projections about nutrient responses under differing conditions must be made cautiously, as soil chemistry appears to play a major role in their expression. Our results suggest that no single nutrient optimum exists, and attempts to identify a reasonable balance among plant nutrients cannot be successful using foliar or fruit analysis alone or using the visual appearance of plants. Clearly, the study of nutrient interactions will remain a challenge to horticulturists for many years to come.

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On 23 Apr. 1987, certified nursery plants of ‘Earlglow’ were planted 30 cm on square, forming a 40-plant grid in each plot. Standard cultural practices were followed (Pritts et al., 1988), with the exception that runners were removed so plants did not form a solid bed. Napropamide at 2.2 kg·ha⁻¹ was applied 2 weeks after planting and again at 4.5 kg·ha⁻¹ before straw mulch application in November. Ammonium nitrate was applied 6 weeks after planting to supply 34 kg N/ha, and calcium nitrate was applied in early September to supply 22 kg N/ha. Soil moisture was maintained near field capacity throughout the study with overhead irrigation, as indicated by soil moisture tensiometers.

Runners were removed weekly to maintain a constant plant density within each plot. Entire plants were excavated beginning 2 June 1987 and continuing at 30-day intervals throughout the first year of establishment (1987) and first fruiting season (1988). Plants were separated into roots, crowns, petioles, leaf blades, and fruit, washed with tap water, then rinsed with distilled water. Plant material was oven-dried at 65°C, weighed, and ground in a Wiley mill to pass through square grids 1.4 × 1.7 mm.

Nitrogen concentration in leaf lamina was determined by a Kjeldahl procedure. The P, K, Ca, Mg, Fe, Zn, B, Mn, Cu, and Mo content was determined by inductively coupled plasma spectroscopy (Model 975 Plasma Atomcomp ICP Spectrograph with ICAP 61 Update, Jarrell-Ash, Pittsburgh). Leaf area was determined using a portable area meter (LI-COR Model LI-3100, Lincoln, Neb. Aboveground plant dry weight, number of branch crowns and fruit per inflorescence, inflorescence length, fruit count per plant, fruit weight, achenes per square centimeter of surface, and fruit weight per achene were recorded at each sampling date. Yield component data were pooled for each plot. A final leaf sample for tissue nutrient analysis was collected ≈5 weeks after postharvest leaf removal. Fruit soluble solids concentration (SSC) was determined with a hand-held refractometer (Thomas Scientific, Swedesboro, N.J.).

Statistical analysis of the central composite design was used to derive meaningful regression equations that accurately described the response surfaces for the variables of interest. Values for nutrient concentrations were transformed before analysis by first subtracting the mean, then dividing by it, resulting in a scale from -1 to 1 for each nutrient. Using a transformed scale in regression analysis has three advantages over using the actual levels of applied nutrient. First, regardless of the nutrient involved, -1 represents a low level and +1 a high level. This scheme is important when the differences in ranges between low and high levels are very large (as was the case with P and B). Second, the transformed variables are orthogonal, allowing unbiased estimates of regression coefficients to be calculated. Third, regression coefficients among components of the model can be compared directly with each other. Also, this standardized scale can be converted easily to the actual applied values through multiplication.

These new, orthogonal variables were used in regression analysis to calculate the response surface, using the following full model: Response = Block + P + Zn + B + P × Zn + P × B + B × Zn + P² + B² + Zn². Sequential sums of squares were accumulated for the full model. Single-degree-of-freedom F tests were then used to evaluate the significance of the individual components of the model. For the generation of response surfaces, new models were constructed using only the significant terms from the full model, with the exception of fruit count per inflorescence, where the full model was used (since the full model was significant, but individual coefficients were not).

**Results**

Preplant, soil-applied nutrient additions of P, Zn, and B were correlated with soil test results in each of the 2 years (Table 2). A decrease of ≈0.5 pH units was measured during the course of the experiment in plots that initially received lime applications. Plant growth variables were unaffected by liming; however, yield components were affected differently at pH 5.5 and 6.5, so results are discussed separately.

*Results at pH 6.5.* Yield increased with increasing B at a high P level but decreased at a low P level (Table 1, Fig. 1 top).

Table 2. Significant (P < 0.01) correlation coefficients (r) between mineral nutrient treatment levels (1986) and soil test values (1987 and 1988).

| Soil test | P | Zn | B |
|-----------|---|----|---|
| Spring 1987 | 0.94 | 0.97 | 0.84 |
| Fall 1987 | 0.74 | 0.65 | **NS** |
| Spring 1988 | 0.55 | 0.93 | 0.71 |

**NS**Non-significant.
with yield. For K, P, Ca, Mn, Cu, and Zn, foliar and fruit levels were correlated positively, but no correlation existed for levels of Mg, Fe, or B. Plant size was not correlated with yield.

Results at pH 5.5. Strawberry yields were significantly affected by B (Fig. 6), but the interaction of B with P or Zn was not significant (Table 3). Boron applications increased yield regardless of P and Zn levels. As at the higher pH, fruit count per inflorescence \( (r = 0.65, P < 0.001) \) and individual fruit weight \( (r = 0.31, P < 0.04) \) were correlated with yield, and their responses to nutrient additions somewhat paralleled the yield response (Figs. 7 and 8). Again, although the response of fruit count per inflorescence was very similar to that of yield, regression coefficients that describe the response of this variable to nutrient additions were not significant (Fig. 8).

Nutrient additions had no effect on inflorescence count, fruit SSC, or achene spacing (Table 3). Inflorescence length, however, was positively correlated with fruit size \( (r = 0.39, P < 0.009) \), yield \( (r = 0.36, P < 0.02) \), achene spacing \( (r = 0.34, P < 0.02) \), and the level of applied B \( (r = 0.31, P < 0.04) \).

Leaf P at renovation showed a quadratic response to soil-applied P (data not shown), similar to that at the higher pH (Fig. 5). Leaf B exhibited a quadratic response to P \( (P < 0.06) \) but not to B additions (data not shown).

Leaf Zn at renovation was negatively correlated with yield \( (P < 0.06) \), but correlations of yield with other foliar nutrient levels at renovation were not significant (data not shown). Fruit and leaf nutrient levels were correlated positively for P, Mn,

Fig. 3. Response of branch crowns per plant to P and B at Zn = 0 (23.9 kg·ha\(^{-1}\)) and pH = 6.5. Branch crown count = 11.0 – 3.15(P), \( R^2 = 0.09, P < 0.05 \). For scale conversion, units of P range from 11.2 (–1) to 123.3 (1) kg·ha\(^{-1}\) and B from 0.2 (–1) to 4.5 (1) kg·ha\(^{-1}\).

Fig. 4. Seasonal pattern of P, B, and Zn concentration in leaf lamina of plants set in Apr. 1987 and harvested over the following 16 months for the central treatment of the design of the experiment (kg·ha\(^{-1}\): 67 P, 23.9 Zn, and 2.35 B).

Fig. 5. Response of leaf P at renovation to P and B at Zn = 0 (23.9 kg·ha\(^{-1}\)) and pH = 6.5. Percent leaf P = 0.201 + 0.014(P) + 0.034(P\(^2\)) – 0.013(B) – 0.03(B\(^2\)Zn), \( R^2 = 0.339, P < 0.002 \). For scale conversion, units of P range from 11.2 (–1) to 123.3 (1) kg·ha\(^{-1}\) and B from 0.2 (–1) to 4.5 (1) kg·ha\(^{-1}\).

Fig. 6. Yield response (grams per six plants) to P and B at Zn = 0 (23.9 kg·ha\(^{-1}\)) and pH = 5.5. Yield = 1427 + 410 (P) + 166(B), \( R^2 = 0.153, P < 0.03 \). For scale conversion, units of P range from 11.2 (–1) to 123.3 (1) kg·ha\(^{-1}\) and B from 0.2 (–1) to 4.5 (1) kg·ha\(^{-1}\).
but we did find a yield response to B that was mediated both through fruit count per inflorescence and fruit size. Although the response of fruit count per inflorescence to B was not statistically significant, this yield component was highly correlated with yield at both pH levels, and the response surfaces were similar (Figs. 1, 2, 6, and 7).

Boron influences auxin activity (Pilbeam and Kirkby, 1983), and since auxin is related to both inflorescence development and fruit receptacle expansion (Archbold and Dennis, 1985), one would expect yield to respond to supplemental B if its ambient levels were inadequate and P was adequate. Inflorescence length may be a useful indicator of auxin activity, as this variable was associated with achene spacing on the receptacle and was positively associated with yield, fruit size, and applied B at the lower pH.

Interactions among nutrients may help explain inconsistent responses to applied nutrients that have been reported in the literature. Neilson and Eaton (1983) studied the effects of B nutrition on strawberry yield components using solution culture and found no yield response to B applications from 0.01 to 1.25 ppm. Riggs et al. (1987) found that B applications up to 4.5 kg·ha⁻¹ decreased yields of ‘Tritic’ and ‘Benton’ strawberries on Quatama loam soils suspected to be low in B in the Willamette Valley, Ore. In our study, at a soil pH of 5.5, yield responded positively to B. However, at a soil pH of 6.5, the responses to applications of B were either positive, neutral, or negative, depending on the specific level of P fertilization (Fig. 1).

Our results demonstrated an important shortcoming of field studies that examine single nutrients, without considering interactions with other nutrients. Adams (1980), Kirsch (1959), Murphy et al. (1981), and Sumner and Farina (1986) emphasized the importance of designing nutritional experiments that permit measurements of potential interactions to avoid possible misleading main-effect conclusions.

While it is important for horticulturists to attempt to identify the nutritional conditions that optimize total yield, it is likely that all of the various components contributing to final yield do not have the same nutritional requirements. For example, number of branch crowns, fruit count per inflorescence, fruit weight, and runner production responded differently to additions of P, Zn, and B. Also, responses changed with soil pH. These observations and the finding of interactions suggest that there may not be a single optimum combination of nutrients that maximizes yield. Yield optimization, in reality, likely consists of multiple sets or nutrient levels or ranges, which of themselves are not limiting to plant growth or yield.

There are many inconsistencies in the literature describing the response of strawberries to fertilization (May and Pritts, 1990). Some of these inconsistent responses may reflect nutrient interactions such as those found in our study, while others may reflect variations in climate, soil moisture, soil type, plant condition, and cultural practices. Tissue analysis has been viewed as an invaluable tool to assess the nutritional status of perennial crops and to directly evaluate plant response across a wide range of conditions (Bould, 1964). However, in our study, leaf nutrient levels during the recommended postharvest period in 1988 generally were not related to yield, nor were foliar nutrient concentrations in 1988 correlated with nutrient levels in the soil.

At pH 6.5, applied P was correlated with leaf P, which in turn was correlated with fruit P. Applied Zn was not correlated with foliar Zn, although foliar and fruit Zn levels were correlated. No correlation existed among soil, foliar, or fruit B or between yield and fruit P, B, or Zn. Further, B levels in ripening fruit (17 to 22 ppm) were much lower than foliar levels at the same time (54 to 73 ppm). Therefore, practices that increase soil nutrient levels may not lead to increased foliar levels, especially if nutrient dilution occurs as growth is stimulated. In addition, a strategy that successfully increases foliar nutrients may not always lead to higher levels in reproductive tissues. Finally, higher levels in reproductive tissues may not be associated with higher yield or fruit quality.

Others have concluded that foliar analyses are not particularly useful indicators of nutrient status for particular elements. Riggs et al. (1987) found that soil applications of B were not effective in increasing total B in reproductive tissues and that foliar B levels were a poor predictor of levels in reproductive tissues.

A plant nutrient analysis alone may not be sufficient to identify treatments that will increase yield economically, at least when plants are grown on a reasonably fertile soil. The challenge of understanding nutritional responses is great, considering the thousands of possible interactions that could occur among the 12 mineral nutrients for a single variable under a single set of conditions. Adding to the complexity of evaluating plant nutrient status is the observation that leaf nutrient levels fluctuate seasonally (Fig. 4). Even if one could find specific times when the level of certain nutrients is predictive of plant performance, growers may not accept the time and expense involved with multiple sampling times.

Finally, the results of our study are limited because we examined the response of only a single genotype. Some growers have reported that ‘Earliglow’ is relatively tolerant to low nutrient levels, but this evidence is anecdotal. With many crop plants, including strawberries (John et al., 1976; Peterson et al., 1986; Renquist and Hughes, 1985), significant genetic variation exists in the ability to extract or absorb nutrients from the rhizosphere or tolerate low levels of nutrients. More work will be required to identify genotypes tolerant to nutrient imbalances.

Nutrient additions significantly affected strawberry yield through various yield components, and these responses were pH dependent. Interactions among applied nutrients also occurred, making it difficult to predict strawberry plant response under different conditions. Soil type, climate, cultivar, plant condition, cultural practices, and the levels of other plant nutrients also likely influence the response of the strawberry plant to P, B, and Zn. Projections about nutrient responses under differing conditions must be made cautiously, as soil chemistry appears to play a major role in their expression. Our results suggest that no single nutrient optimum exists, and attempts to identify a reasonable balance among plant nutrients cannot be successful using foliar or fruit analysis alone or using the visual appearance of plants. Clearly, the study of nutrient interactions will remain a challenge to horticulturists for many years to come.

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