

NITROGEN AND POTASSIUM FERTILIZATION OF POTATOES:
EVALUATING NUTRIENT ELEMENT INTERACTIONS IN PETIOLES
WITH RESPONSE SURFACES¹

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Abstract

We studied the effects of fertilizer N and K rates on the nutrient concentrations in petioles of Russet Burbank potato grown on a low-K, low-Cl, highly calcareous Millville silt loam soil. Phosphate and CaSO_4 were applied uniformly so that P and S did not limit plant growth. An incomplete factorial experimental design, utilizing 14 combinations of fertilizer N and K, including two forms of K, was used to develop a multi-dimensional regression model. This model was then used to produce response surfaces as an aid to analysis of the results and to illustrate the fertilizer treatment effects and their interactions on petiole chemistries. Nitrogen, K and Cl fertilizers had complex effects on petiole concentrations of $\text{NO}_3\text{-N}$, $\text{SO}_4\text{-S}$, Cl, K, Ca and Mg. Nitrate-N and Cl were mutually antagonistic. Soil N had a large positive effect on petiole K and Mg levels and, to a lesser extent, petiole Ca levels. There was competition between K and Mg in petiole concentration. The sum of K, Ca and Mg was essentially constant across fertilizer treatments and sampling times. The sum of $\text{NO}_3\text{-N}$, $\text{SO}_4\text{-S}$ and Cl was highly variable. The ratio of cations to anions ranged between three and five across fertilizer treatments and sampling times. The data indicate that calibration of the major nutrient element concentrations in potato petioles, as a guide to fertilization, requires knowledge of background soil fertility conditions with respect to N, K, Cl, Ca and Mg, and an understanding of the antagonisms and synergisms among these elements.

Compendio

Se estudiaron los efectos de las dosis de fertilizantes nitrogenados y potásicos sobre las concentraciones en los peciolos de papa Russet Burbank creciendo en suelo Millville aluvión-franco, altamente calcáreo, con bajos contenidos de K y Cl. Se aplicó uniformemente fosfato y SO_4Ca de manera que el P y el S no limitaran el crecimiento de las plantas. El diseño experimental fue un factorial incompleto, utilizando 14 combinaciones de fertilizantes N y K, incluyendo dos formas de K, para desarrollar un modelo

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de regresión de dimensión múltiple. Este modelo fue luego utilizado para producir superficies de respuesta como un medio para analizar los resultados e ilustrar los efectos del tratamiento con fertilizantes y sus interacciones sobre las composiciones químicas de los peciolo. Los fertilizantes nitrogenados, potásicos y clorados (N, K y Cl) tuvieron efectos complejos sobre las concentraciones de $\text{NO}_3\text{-N}$, $\text{SO}_4\text{-S}$, Cl, K, Ca y Mg. Los nitratos y el Cl fueron mutuamente antagonísticos. El nitrógeno del suelo tuvo un efecto positivo prolongado sobre los niveles de K y Mg en los peciolo y en menor grado sobre los niveles de Ca. Hubo competencia entre las concentraciones de K y Mg en el peciolo. La suma de K, Ca y Mg fue esencialmente constante en todos los tratamientos con fertilizantes y momentos de toma muestras. La suma de $\text{NO}_3\text{-N}$, $\text{SO}_4\text{-S}$ y Cl fue muy variable. La relación de cationes a aniones varió entre tres y cinco en los tratamientos con fertilizantes y momentos de la toma de muestras. Los datos indican que la calibración de las concentraciones de los principales elementos nutrientes en los peciolo de papa, como una guía para la fertilización, requiere el conocimiento de los antecedentes de las condiciones de fertilidad del suelo con respecto a N, K, Cl, Ca y Mg y de los antagonismos y sinergismos entre estos elementos.

Introduction

The nutrient concentration of plant tissues, together with diagnostic soil tests, are used to help achieve least cost per unit of crop production. Plant analysis is based on the principle that the concentration of nutrients within the plant, or one of its parts, integrates all of the factors affecting plant growth, including the availability of elements in the soil (15). Plant analysis as an aid to fertilization of vegetable crops, including potato, was discussed by Geraldson and Tyler (5). They summarized the nutrient concentrations associated with deficiency and sufficiency and the relationship of plant growth stage, cultivar, soil fertilization and nutrient interactions in relation to optimal levels of nutrients. Genetic potential (cultivar) governs nutrient concentrations in plants as well as crop yield potential (9).

Potassium chloride is an economical source of fertilizer K but Cl may confound K soil-plant nutrient relationships. Chloride has three different plant functions: (a) as a plant nutrient; (b) through competition with other nutrients in plant up-take; and (c) as a component of the cellular osmotic. Chloride is not a frequent subject of field soil fertility investigations, but Cl was included here because the soil and irrigation water at the experimental site were very low in this element.

Chloride is an essential nutrient element for various higher plants (1). Chloride deficiencies have been reported for potato (4), red clover (22) and kiwifruit (18), all of which were done under greenhouse conditions.

Chloride deficiency in field-grown plants is rare even though other functions of Cl in plants are quite general. For example, there is a striking interaction (mutual antagonism) between Cl and NO_3 for plant uptake in

potato (7); barley (3); sugarbeet and fodderbeet (8, 14); bean and snapbean (23); kiwifruit (18); and tomato (10).

Interactions among inorganic anions (Cl^- , SO_4^{2-} , NO_3^- , H_2PO_4^-) occur both at the soil-root interface and within the plant cells (6). Hiatt and Leggett (6) state also that endogenous interactions include organic anions and a feedback mechanism that influences nutrient element uptake by roots.

Whereas Cl is required in small amounts by plants, under some conditions it may accumulate in large amounts in plants (2) and Cl toxicity may occur in sensitive crops such as soybean, even under non-saline conditions (17). Chloride uptake lessens the production of organic anions by the plant (2). It is not known whether there is a savings in metabolic energy when Cl replaces some organic anions.

One of the primary roles of K in the plant is as an osmoticum and normally Cl acts as a non-metabolized counter-ion to K (2). It is apparent therefore, that the roles of Cl in the plant are strongly affected by other elements. Nevertheless, Flowers (2) emphasized that Cl has a unique role in plant nutrition since a deficiency results in visual symptoms and growth retardation.

The research reported here sought to determine the optimum N and K soil fertility conditions for production of Russet Burbank potato growing on a low-K, calcareous soil. The Cl- NO_3^- interactions, the K-Ca-Mg interactions, and the cation and anion balance in the petioles of Russet Burbank potatoes are described. Other papers in this series examine the fertilizer treatment effects on tuber yield and specific gravity (20) and the effects on tuber sugars and starch (21).

TABLE 1.—*Experimental design; fourteen treatment combinations in two incomplete NxK factorials based on K source.*^a

K		kg N/ha			
		0	112	224	336
Source	kg/ha				
		-----Treatment number-----			
Cl	0	1		2	
	112		3		4
	224	5		6	
	336				
	448		7		8

SO ₄	0	1		2	
	112		9		10
	224	11		12	
	336				
	448		13		14

^aTreatments 1 and 2 were repeated in the Cl and SO₄ subsets for generating the respective response surfaces. N was supplied as urea. K was supplied respectively as KCl and K₂SO₄. Design consists of two interlocking 2x2 factorials.

Materials and Methods

Soil and Crop

Russett Burbank potatoes were grown near Logan, Utah, on a Millville silt loam (coarse silty carbonatic mesic typic Haploxerolls), a well drained, strongly calcareous soil formed from dolomitic parent material. Characteristics of the soil and irrigation water at the site are reported by Westermann *et al.* (20). Briefly, this soil was low in K fertility and the irrigation water was very low in dissolved K, Cl and SO_4 .

Experimental Design

These experiments were specially designed to produce response surfaces of the dependent variables (petiole nutrient element concentrations in this paper) to aid in evaluation of the interactions among potato nutritional factors. Fertilizer treatment combinations and other features of the experimental design are given in Table 1: Fourteen N and K fertilizer treatment combinations were arranged in an incomplete factorial design with four replications. The experimental design is described in more detail elsewhere (20). Phosphorus as 0-45-0 (50 kg P ha⁻¹) and S as gypsum (30 kg S ha⁻¹) were applied uniformly to avoid deficiencies of these elements in the crop.

Petiole Analysis

Petiole samples were collected 61, 79 and 106 days after planting in 1988 and 74 and 101 days after planting in 1989. About 30 petioles, the fourth leaf from the growing tip, were collected from the middle two rows of each plot.

The petioles were dried at 60 C, and ground to pass a 40 mesh screen. The petiole tissue was analyzed for $\text{NO}_3\text{-N}$ by a specific-ion electrode (13).

TABLE 2.—Regression model used to produce response surfaces of the dependent variables.^a

Regression Coefficient	Main and interaction effects
b0	overall mean
b1	N
b2	K
b3	Cl-SO ₄ (a)
b4	N ²
b5	K ²
b6	NxK
b7	NxCl-SO ₄
b8	KxCl-SO ₄
b9	N ² xCl-SO ₄
b10	K ² xCl-SO ₄
b11	NxKxCl-SO ₄

^aRead Cl-SO₄ as Cl or SO₄. These were represented in the model as categorical variables, *i.e.* (+1) and (-1) respectively.

The tissue was dry ashed at 500 C for 6 h and the residue dissolved in 50 ml of 0.2 N HNO₃. This digest was analyzed for P (11); K, Ca, and Mg (atomic absorption spectrophotometry); and Cl and S by flow injection analysis (12; Method Nos. 12-117-07-1-A and 12-116-10-1-C, respectively.) Nitrate-N is reported as mg/g (parts per thousand) dry weight. All other elements are reported as percent dry weight. For some comparisons the petiole nutrient concentrations were converted to millimoles charge per gram (mmol_c/g) and from this the sum of cations, sum of anions and their ratios were computed.

Statistical Analyses

Preliminary two-way analyses of variance were calculated on the potato petiole chemistry parameters. Where there were significant treatment effects, the treatment means were used to generate a regression equation based on the first order regression model of Table 2. The regression equation was used to interpolate the missing cells in the incomplete factorials (Table 1). In other words, the 14 treatment combinations were used to generate 20 estimated treatment means in each of the N-KCl and N-K₂SO₄ response surfaces.

Only a portion of the response surfaces arising from the experiments are presented in the results to conserve space. Westermann *et al.* (20) emphasize that even though there was considerable range in N and K fertilizer rates, the expected optimum N and K combinations were located under the central region of the tuber yield response surface.

The "goodness of fit", *i.e.* R², between the measured treatment means and their counterparts predicted by the model, was computed.

Results and Discussion

Table 3 summarizes the analysis of variance for the fertilizer N, KCl and K₂SO₄ treatment main effects on individual nutrient element concentrations

TABLE 3.—Analysis of variance: Treatment main effects on petiole nutrient element concentrations.

Dependent variable	Sampling time-Year				
	1-88	2-88	3-88	1-89	2-89
	F ratio*				
NO ₃	32.45	27.90	27.99	140.41	57.57
Cl	51.13	25.36	31.54	103.39	6.46
S	2.42	10.77	11.36	8.47	8.60
K	23.71	24.25	14.76	7.27	7.16
Ca	2.18	2.99	6.61	0.85	5.32
Mg	8.09	5.49	9.28	3.83	2.73
ΣCations	8.63	4.16	4.38	0.85	2.97
ΣAnions	19.60	18.96	10.78	48.15	6.99
ΣCations/ΣAnions	20.15	21.27	10.51	20.46	1.81

*F₀₁ = 3.42; F₍₀₅₎ = 2.34.

TABLE 4.—Multiple correlation coefficients for petiole chemistries; comparing predicted treatment means with observed treatment means.

Petiole parameter	Days after planting/year				
	61/88	79/88	106/88	74/89	101/89
	R^2				
NO ₃ -N	0.985	0.996	0.951	0.980	0.968
Cl	0.985	0.999	0.959	0.988	0.986
K	0.965	0.972	0.983	0.929	0.943
Ca	0.806	0.984	0.956	0.615	0.933
Mg	0.979	0.983	0.976	0.857	0.979
SO ₄ -S	0.812	0.945	0.981	0.975	0.945
ΣCations	0.949	0.958	0.981	0.962	0.908
ΣAnions	0.982	0.993	0.981	0.962	0.908
ΣCations/ΣAnions	0.963	0.984	0.946	0.911	0.849

*All R^2 significant at $P \leq 0.01$.

for each sampling time. As indicated, treatment effects were generally significant; some exceptions were Ca, ΣCations, and ΣCations/ΣAnions in 1989.

TABLE 5.—Comparison of predicted (PR), observed (OB) and standard error of the estimated mean (SE) for selected petiole nutrient element concentrations. First sampling, 1988.

kg K/ha	NO ₃ -N, PPT			
	0	112	224	336
0	PR 9.67 (0.65) OB 9.85	16.91 (0.58)	20.18 (0.60) 19.40	19.47 (1.05)
112	PR 8.47 (0.53) OB	15.86 (0.48) 16.82	19.29 (0.45)	18.74 (0.66) 19.35
224	PR 7.68 (0.62) OB 7.05	15.23 (0.53)	18.80 (0.56) 18.25	18.40 (0.60)
336	PR 7.30 (0.83)	15.00 (0.49)	18.73 (0.54)	18.48 (0.60)
448	PR 7.32 (1.29) OB	15.18 (0.68) 15.55	19.06 (0.58)	18.97 (0.69) 18.80
kg K/ha	K %			
	0	112	224	336
0	PR 5.46 (0.16) OB 5.65	5.29 (.14)	5.06 (0.14) 4.78	4.76 (0.25)
112	PR 6.51 (0.12) OB	6.48 (0.12) 6.52	6.38 (0.11)	6.22 (0.16) 6.42
224	PR 7.13 (0.15) OB 6.88	7.243 (0.13)	7.28 (0.13) 7.35	7.26 (0.14)
336	PR 7.33 (0.20)	7.58 (0.12)	7.75 (0.13)	7.87 (0.14)
448	PR 7.10 (0.31) OB	7.49 (0.16) 7.65	7.80 (0.14)	8.06 (0.16) 7.92

kg K/ha	Cl%			
	0	112	224	336
0	0.74 (0.09) OB 0.62	0.14 (0.08)	-0.05 (0.08) 0.11	0.18 (0.14)
112	PR 1.67 (0.07) OB	0.92 (0.06) 0.94	0.58 (0.06)	0.65 (0.09) 0.54
224	PR 2.41 (0.08) OB 2.56	1.50 (0.07)	1.02 (0.07) 0.93	0.94 (0.08)
336	PR 2.96 (0.11)	1.91 (0.06)	1.27 (0.07)	1.04 (0.08)
448	PR 3.34 (0.179) OB	2.13 (0.09) 2.04	1.34 (0.08)	0.96 (0.09) 1.05

kg K/ha	Ion Ratio			
	0	112	224	336
0	3.24 (0.09) OB 3.34	2.58 (0.08)	2.22 (0.08) 2.16	2.16 (0.14)
112	PR 2.96 (0.07) OB	2.38 (0.06) 2.24	2.09 (0.06)	2.09 (0.09) 2.13
224	PR 2.73 (0.08) OB 2.65	2.21 (0.07)	2.00 (0.07) 2.14	2.07 (0.08)
336	PR 2.53 (0.11)	2.09 (0.06)	1.94 (0.07)	2.09 (0.08)
448	PR 2.38 (0.17) OB	2.01 (0.09) 2.07	1.94 (0.08)	2.15 (0.09) 2.09

The regression model predicted nutrient element concentrations for all N and K rate and source combinations in the complete 4N x 5K complete factorial (Table 1). The "goodness of fit" (R^2) for each sampling time is shown in Table 4. "Goodness of fit" ranged from 0.615 to 0.988, averaging 0.946 across all sampling times and years. The model was also efficient in estimating tuber yields and specific gravities; R^2 ranged between 0.72 and 0.98 for these parameters (20). The predicted treatment mean, the observed treatment mean and the standard error of the estimate for selected parameters in 1988 are presented in Table 5. For economy of space other parameters and sampling times in 1988 and 1989 are not given but the response surface-generating model was equally reliable for all other data sets.

The following figures illustrate the many treatment effects and interactions that were encountered. Response surfaces omitted because of space were equally significant and supportive of the final conclusions. The surface of each three-dimensional graph is rotated so that it slopes downward to the viewer to avoid concealing lines. Accordingly, the orientation of each figure may vary to provide the clearest view of each data set, as does the scaling of the axes. It will be noted that conformation of the response surface changes from element to element and among sampling times. In this manner of presentation the individual treatment means are less important than are the relationships among means, *i.e.* interactions.

Figure 1 shows the treatment effects on petiole- $\text{NO}_3\text{-N}$. The base level of petiole- NO_3 was higher in 1988 than in 1989 (Fig. 1a vs 1b), partially

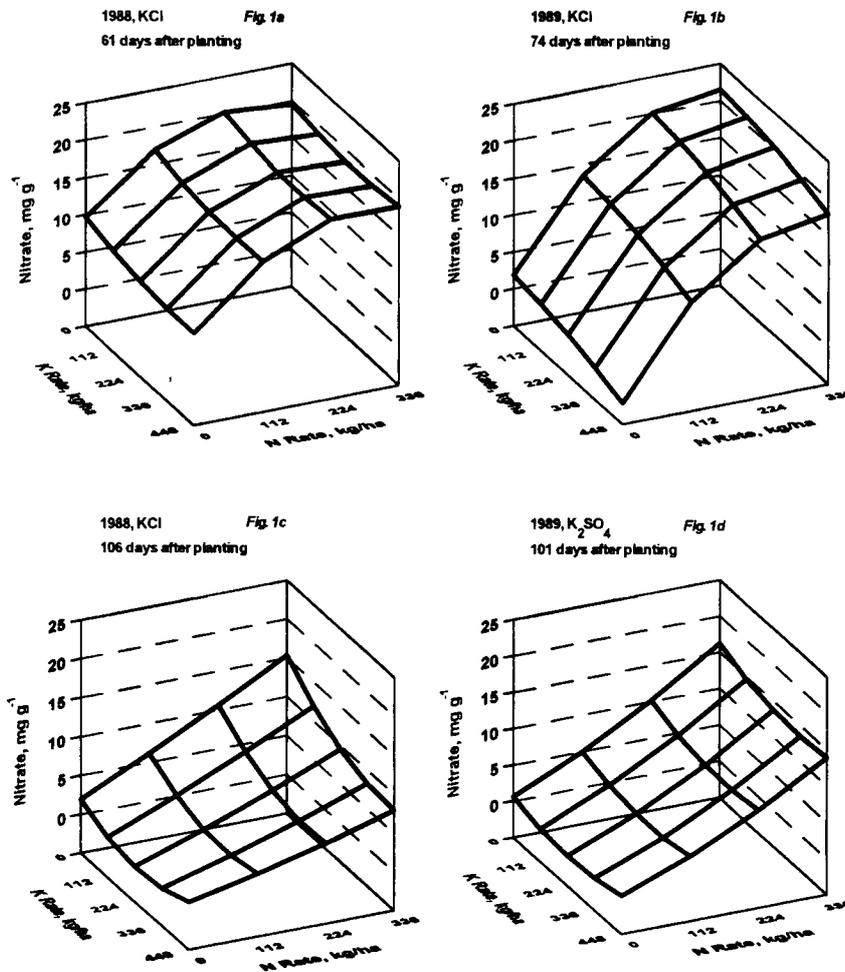


FIG. 1. Petiole $\text{NO}_3\text{-N}$ responses to N and KCl (a, b, c) and K_2SO_4 (d) fertilizers at selected sampling times in 1988 and 1989.

because the first sampling was earlier in 1988. Petiole- NO_3 decreased rapidly with time (Fig. 1a vs 1c). In both seasons petiole- NO_3 increased with fertilizer-N to a maximum of about 20 mg/g at 224 kg N/ha. Fertilization with KCl sharply decreased $\text{NO}_3\text{-N}$ petiole levels after the first sampling. Fertilization with K_2SO_4 did not significantly decrease petiole NO_3 levels.

Initially the fertilizer-N was urea, but nitrification converts the bulk of the fertilizer-N to NO_3 in Millville silt loam soil within about three weeks after application. Accordingly, the plant would be exposed to essentially 100% of the fertilizer-N as NO_3 for the most of the growing season.

Petiole-Cl concentrations increased with the fertilizer-KCl rate and decreased by 45% to 85% with fertilizer-N in both seasons, depending on the soil treatment (Figs. 2 a and c). Whereas $\text{NO}_3\text{-N}$ generally decreased with

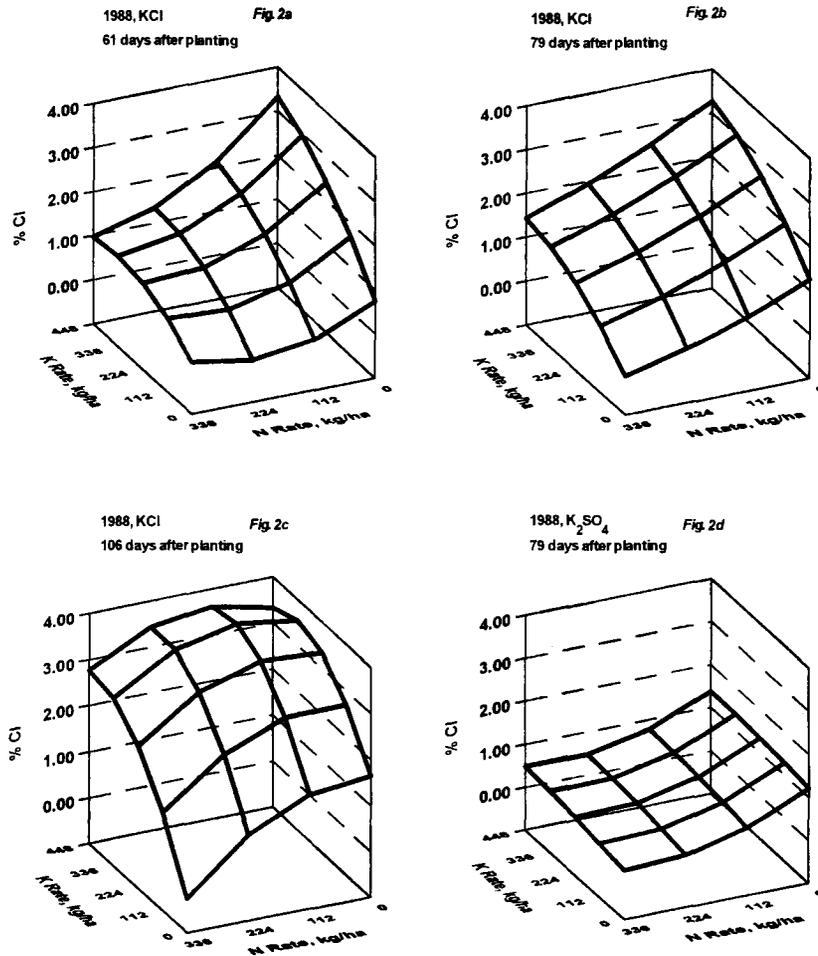


FIG. 2. Petiole Cl response to N and KCl (a, b, c) and K_2SO_4 fertilizers at selected sampling times in 1988.

time (Fig. 1), petiole-Cl increased (Fig. 2). Petiole-Cl tended to increase with fertilizer K_2SO_4 (and decrease with fertilizer-N), probably because of the small amount of Cl present in the K_2SO_4 (Fig. 2d). The overall results in Figs. 1 and 2 show that the NO_3 -Cl antagonism is reciprocal. The large effects of fertilizer-Cl on petiole- NO_3 has been observed elsewhere (7, 10). Unpublished data from Kimberly, Idaho show that the NO_3 -Cl antagonism occurred at relatively high background Cl levels also.

Gausman *et al.* (4) reported that Cl deficiency was seen in potato where leaf tissue contained 0.0143 to 0.0381 mmol Cl/g corresponding to 0.051% and 0.13% Cl dry weight. In our study Cl was as low as 0.068% in individual plots. Petiole-Cl averaged 0.108% where KCl was not applied. It appears that Cl might have been deficient in this low-Cl field environment where it

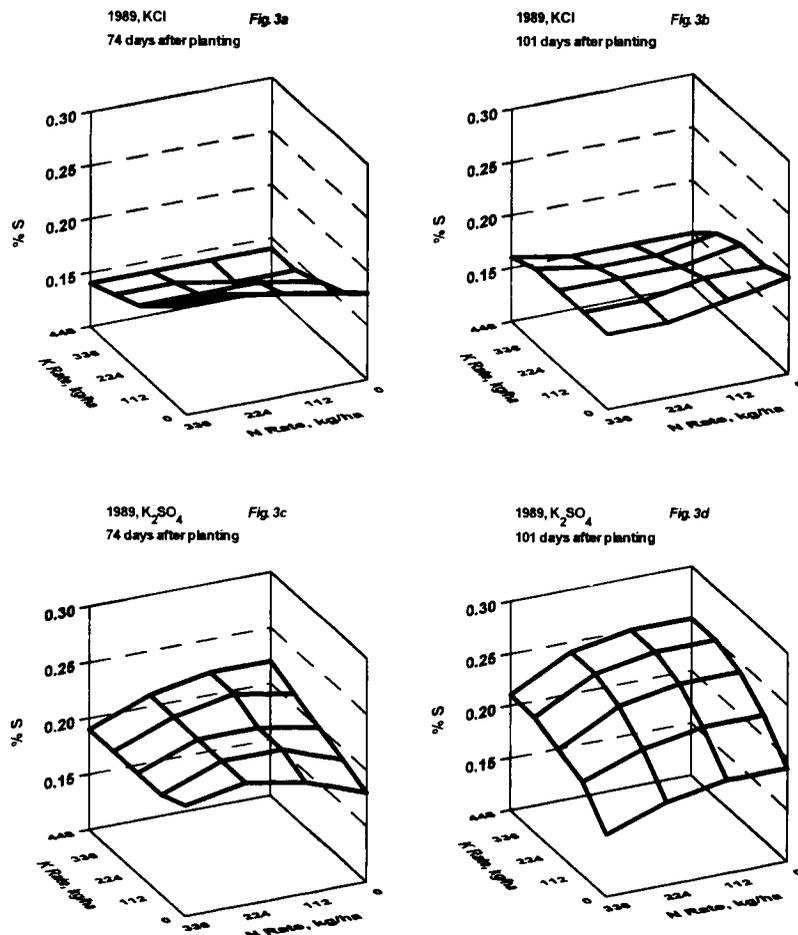


FIG. 3. Petiole $\text{SO}_4\text{-S}$ responses to N and KCl (a, b) and K_2SO_4 (c, d) fertilizers at selected sampling times in 1989.

was not included in the fertilizer.

The $\text{NO}_3\text{-Cl}$ antagonism is of considerable practical consequence. Although it seemed to have little effect on tuber yield, it did affect specific gravity (20) and markedly affected tuber starch, sucrose and reducing sugar (21). Our results indicate that petiole analyses, as a guide to N fertilization of potatoes, should be calibrated against background levels of Cl in the environment.

Fertilizer treatment effects on petiole $\text{SO}_4\text{-S}$ are illustrated in Fig. 3. Petiole- SO_4 tended to decrease with KCl rates but there was little apparent effect of fertilizer-N. By contrast, petiole- SO_4 increased markedly with fertilizer-S and decreased with fertilizer-N (Fig. 3d), possibly because of dilution in the more vigorous plant growth associated with fertilizer-N. The $\text{NO}_3\text{-SO}_4$ competition was clearly not as pronounced as that for $\text{NO}_3\text{-Cl}$. As

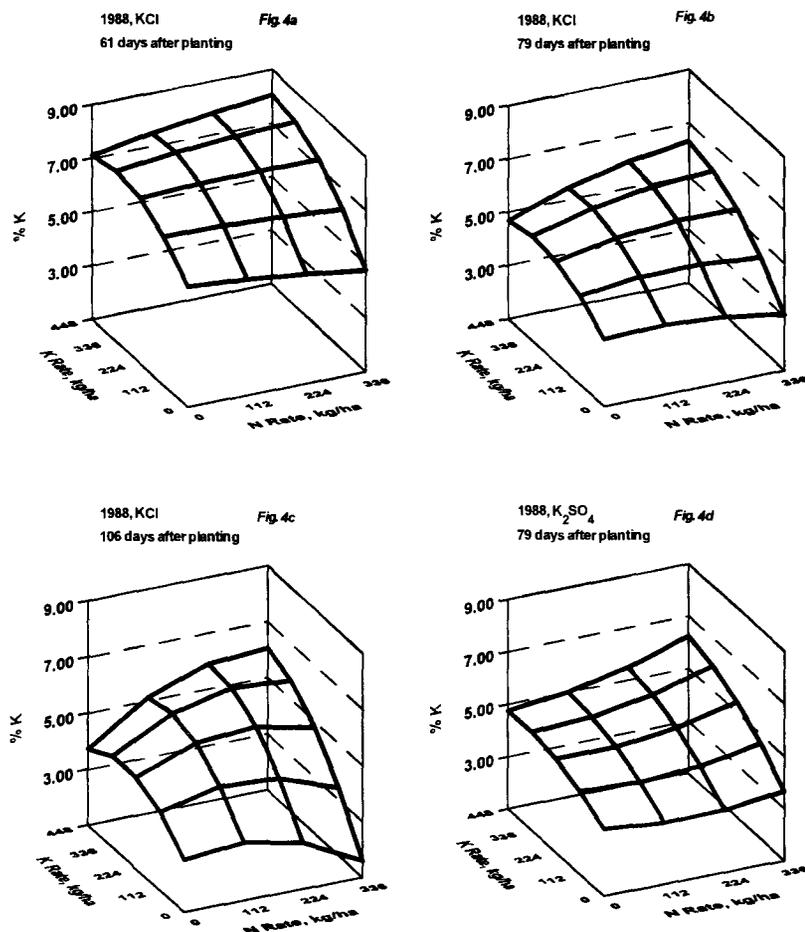


FIG. 4. Petiole K responses to N and KCl (a, b, c) and K_2SO_4 (d) fertilizers at selected sampling times in 1988.

stated above, $CaSO_4$ was applied uniformly to eliminate the possibility of S deficiency. Thus, even though S did not limit plant growth the KCl, K_2SO_4 and N fertilizer treatments collectively had considerable effect on petiole- SO_4 concentrations.

Fertilizer-K increased petiole-K at all petiole sampling times (Fig. 4). Fertilizer-N decreased petiole-K at zero fertilizer-K (probably a dilution effect) but increased petiole-K in the presence of KCl. The positive effect of soil N on plant K became more pronounced over the growing season. With fertilizer K_2SO_4 there was a slight but significant positive effect of fertilizer-N on petiole K (Fig. 4d).

Petiole-Ca concentration increased with sampling time and decreased slightly with fertilizer-K and N (Fig. 5). The treatment effects were more

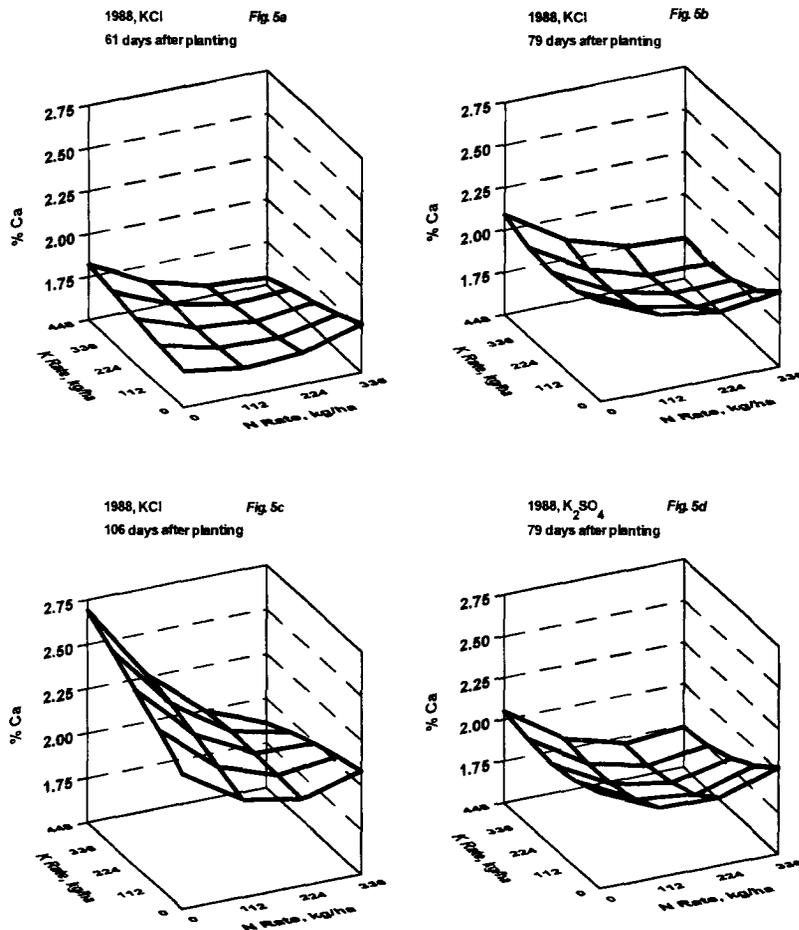


FIG. 5. Petiole Ca responses to N and KCl (a, b, c) and K_2SO_4 (d) fertilizers at selected sampling times in 1988.

significant for the last sampling in 1988 (Table 3). Figure 5c shows that late-sampled petiole-Ca increased sharply with KCl where soil N was low. Thus, the otherwise simple negative interaction between K and Ca was confounded markedly by fertilizer-KCl. The potato plant seemed to balance increased petiole-Cl with petiole-Ca. Potassium sulfate had no effect at zero fertilizer-N on petiole-Ca but at 336 kg N/ha fertilizer-K decreased petiole-Ca (Fig. 5d). Overall, a complex interaction involving cationic and anionic charge balancing is evident. This is discussed further below.

Petiole-Mg increased with sampling time and decreased to some extent with fertilizer-K (Fig. 6). Fertilizer-N tended to increase petiole-Mg at zero fertilizer-K, the tendency decreasing with increasing fertilizer K. The KCl and K_2SO_4 effects were very similar. The effect of KCl on petiole-Ca did not

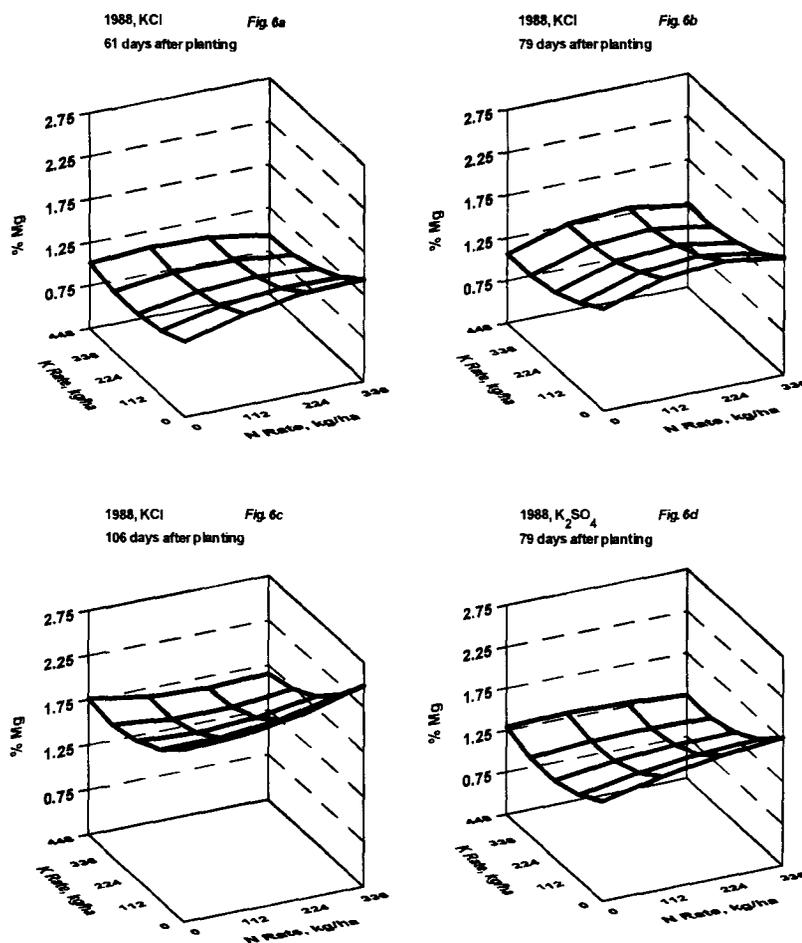


FIG. 6. Petiole Mg responses to N and KCl (a, b, c) and K_2SO_4 (d) fertilizers at selected sampling times in 1988.

occur with petiole-Mg (Fig. 5 vs 6).

Despite the fact that K, Mg and Ca concentrations in petioles varied markedly among fertilizer treatments and over time, the sum of these cations was constant; the response surface was essentially flat for all sampling times (data not shown). By contrast there were highly significant treatment effects on the sum of anions (NO_3 , Cl, SO_4). The summation of nutrient antagonisms and synergisms are given in Fig. 7, which relates the ratio $\Sigma Cations / \Sigma Anions$ to N and K soil treatments. The ion ratio increased with sampling time (Fig. 7). The ratio generally decreased with KCl but the response surface conformation markedly differed with K_2SO_4 (Fig. 7c vs 7d).

The effects of K fertilizer on Mg and Ca concentrations of potato petioles warrant further evaluation to determine if, for example, K fertilization

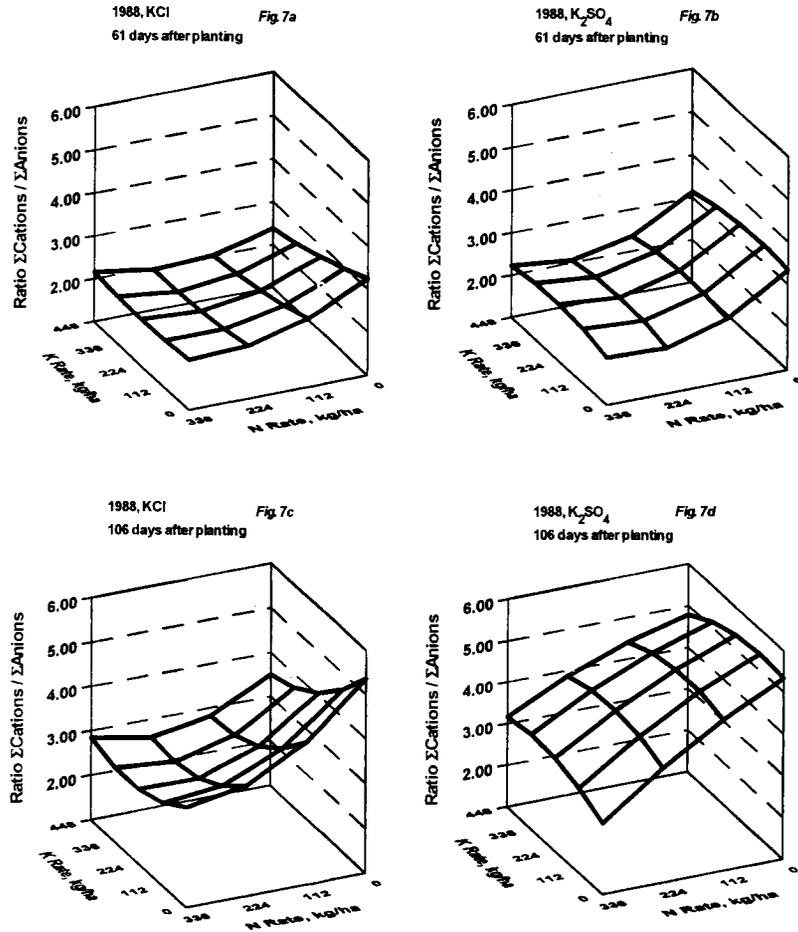


FIG. 7. Petiole $\Sigma\text{Cations}/\Sigma\text{Anions}$ ratio responses to N and KCl (a, c) and K_2SO_4 fertilizers at selected sampling times in 1988.

could induce Mg deficiency in soil derived from dolomite (*i.e.* high levels of available Mg) even though N fertilization induced higher Mg uptake. A study examining the K-Mg-plant relations would need to be carefully designed to segregate the various synergistic and antagonistic interactions on plant nutrient element composition.

The uniformity of $\Sigma\text{Cations}$ across N and K fertilizer levels and time demonstrate (a) that potato adapts readily to general environmental nutrient conditions, and (b) that osmolality of plant cells probably is as important as direct nutritional effects of the individual nutrient elements on plant growth vigor.

Some workers (16, 19) associated increased organic anion production in the plant (*i.e.* $\Sigma\text{Cations} - \Sigma\text{Anions}$) with increased plant yield. In those stud-

ies organic ion production decreased with KCl fertilization in a manner similar to our results. It is apparent that, an exclusive focus on cation-anion balance may ignore the antagonism between NO_3 and Cl and associated changes in the N economy of the plant. Apparent yield differences related to organic anions would be obviated if the NO_3 -Cl antagonism was operating.

The N-K soil treatments did not affect petiole-P concentration, which averaged 0.220% or 0.213 mmol_c/g over the two seasons. Including PO_4 -P in the ΣAnions would uniformly increase ΣAnions by about 16% and decrease the ratio $\Sigma\text{Cations}/\Sigma\text{Anions}$ by about 14%. But with or without petiole-P $\Sigma\text{Cations}$ was three to five times larger than ΣAnions overall. Presumably the difference in ionic electrical charge was balanced by the production of organic acid anions within the plant (16). Evidently Cl decreased the production of organic anions, in agreement with Flowers (2), perhaps because Cl substitutes for organic anions in the plant. Figure 7 shows that fertilizer-S had a similar but somewhat smaller effect.

The lack of petiole-P concentration effects by N, K, S or Cl fertilizer levels, contrast with work of Hiatt and Leggett (6) who reported that plant P concentration depended on other nutrient anions in the rooting medium. Our results might have been different if soil-P was a controlled experimental variable that limited plant growth in some of the treatments. Unpublished data at Logan, Utah show that P fertilization of a low-P soil had a strong positive effect on Mg and Na composition of alfalfa.

Correlations between petiole N and K and tuber yield and specific gravity were described and discussed by Westermann *et al.* (20).

Conclusions

1. Nitrogen, K, and Cl fertilization of Russett Burbank potato grown on a highly calcareous, low-K and low-Cl soil had complex effects on petiole chemistry. The most important of these was the mutual antagonism between NO_3 and Cl, which was clearly demonstrated in this low-Cl environment. The results suggest that petiole- NO_3 is difficult to interpret when the other environmental nutrient conditions are not well characterized.

2. There was a strong positive effect of soil N fertility on K and Mg concentration of potato petioles. This interaction was weak to non-existent for Ca. On the other hand, there was strong competition between K and Mg, and somewhat weak competition between K and Ca in this calcareous soil. Nevertheless, the sum of petiole cation concentrations (expressed as mmol_c/g) was constant across petiole sampling times and soil fertility levels. Evidently, potato has a considerable ability to adjust the osmotica of its vascular system, and probably the osmotica of other organs as well.

3. The sum of inorganic anions in the petiole (*i.e.* NO_3 , Cl SO_4) was highly sensitive to soil N, K and Cl fertility levels. The ratio of inorganic cations to anions in the petiole ranged from 3 to 5, indicating that the

plant produced considerable amounts of organic anions to maintain the ionic charge balance.

4. The incomplete factorial experimental design used in this series of experiments was very efficient in relation to field space and other resources versus the amount of information developed. This statistical tool is particularly useful in evaluating the complex soil fertility-plant nutrient interactions that occur in the field.

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CORRECTION

Figure 1 was inadvertently run twice in the Leever, et al., article in the March (1994) issue of the APJ. Below is the correct Figure 2 to be inserted on page 139.

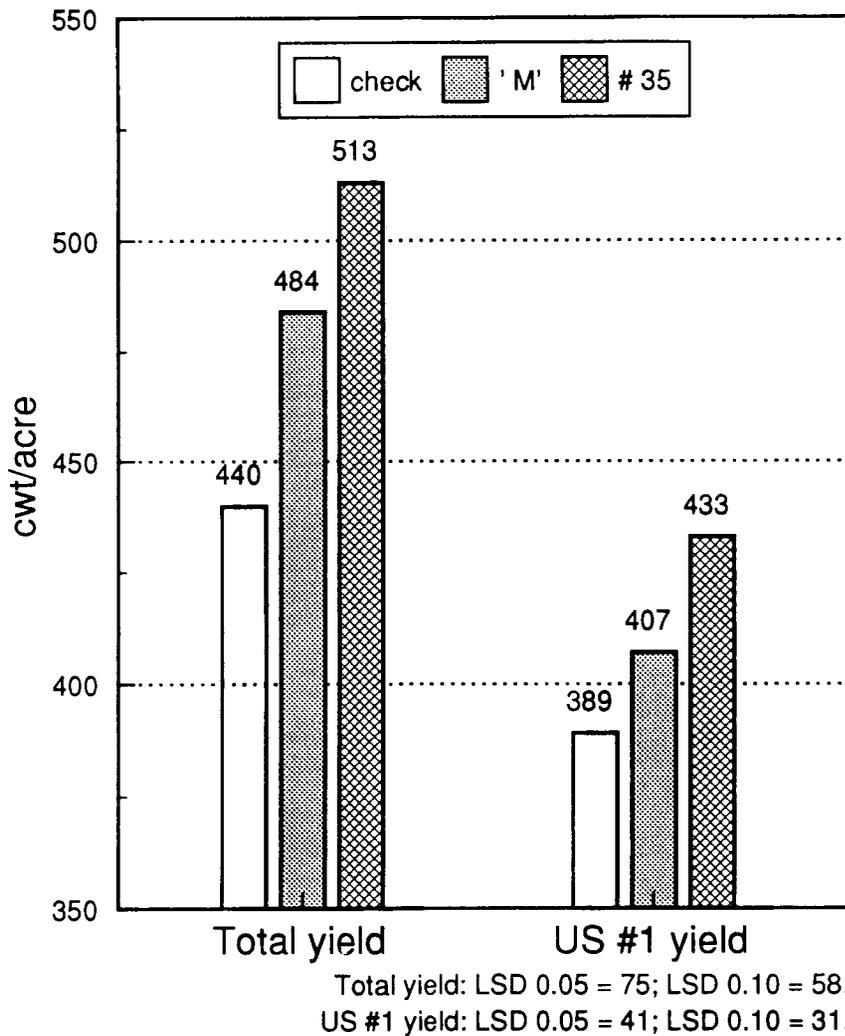


FIG. 2. Total and US#1 yields of Norgold Russet Nebraska strains at Hermiston, Oregon. Excluded is strain #19 (absent in 1 trial).