

NOTA

SPATIAL VARIABILITY OF SOIL CHEMICAL PROPERTIES AFTER COFFEE TREE REMOVAL⁽¹⁾

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ABSTRACT

Assessing the spatial variability of soil chemical properties has become an important aspect of soil management strategies with a view to higher crop yields with minimal environmental degradation. This study was carried out at the Centro Experimental of the Instituto Agronomico, in Campinas, São Paulo, Brazil. The aim was to characterize the spatial variability of chemical properties of a Rhodic Hapludox on a recently bulldozer-cleaned area after over 30 years of coffee cultivation. Soil samples were collected in a 20 x 20 m grid with 36 sampling points across a 1 ha area in the layers 0.0–0.2 and 0.2–0.4 m to measure the following chemical properties: pH, organic matter, K⁺, P, Ca²⁺, Mg²⁺, potential acidity, NH₄-N, and NO₃-N. Descriptive statistics were applied to assess the central tendency and dispersion moments. Geostatistical methods were applied to evaluate and to model the spatial variability of variables by calculating semivariograms and kriging interpolation. Spatial dependence patterns defined by spherical model adjusted semivariograms were made for all cited soil properties. Moderate to strong degrees of spatial dependence were found between 31 and 60 m. It was still possible to map soil spatial variability properties in the layers 0–20 cm and 20–40 cm after plant removal with bulldozers.

Index terms: Geostatistics, soil fertility, traditional management, semivariogram.

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RESUMO: VARIABILIDADE ESPACIAL DOS ATRIBUTOS QUÍMICOS DO SOLO APÓS REMOÇÃO DE CAFEZAL

A avaliação da variabilidade espacial dos atributos químicos do solo tem se tornado importante ferramenta na determinação de estratégias de manejo que visam aumentar a produtividade agrícola com menor degradação ambiental. O presente trabalho foi realizado no Centro Experimental Central do Instituto Agrônomo, localizado em Campinas/SP, com o objetivo de caracterizar a variabilidade espacial dos atributos químicos de um Latossolo Vermelho após a remoção de um cafezal, cultivado por mais de 30 anos, com trator de esteira. As amostras de solo foram coletadas em grade georreferenciada de 20 x 20 m, totalizando 36 pontos nas camadas de 0–0,2 e 0,2–0,4 m, para determinação dos seguintes atributos químicos: pH, matéria orgânica do solo (MOS), K⁺, P, Ca²⁺, Mg²⁺, acidez potencial (H + Al), N-NH₄ e N-NO₃. A variabilidade espacial foi estudada utilizando-se a geoestatística, por meio da análise de semivariogramas e da interpolação por krigagem. Para todos os atributos estudados foi possível estabelecer uma estrutura de dependência espacial definida por um semivariograma ajustado a um modelo esférico. Foram encontradas dependências espaciais de moderada a forte, com valores de alcance variando de 31 a 60 m. Mesmo com a remoção das plantas com trator de esteira, foi possível mapear a variabilidade espacial dos atributos do solo nas camadas de 0–0,2 e 0,2–0,4 m de profundidade.

Termos de indexação: geoestatística, fertilidade do solo, manejo convencional, semivariograma.

INTRODUCTION

Horizontal and vertical variation in the spatial distribution of soil properties results from the interaction of soil formation and management factors (Reichardt et al., 1986; Guimaraes et al., 2001; Cavalcante et al., 2007). Thus, even in pedologically homogeneous areas – with the same geomorphological surface, material of origin and climate – changes in soil use and management may lead to spatial variation in soil properties.

Nowadays, knowledge on the variability of soil properties is considered a basic principle for a localized management in precision agriculture for areas at any scale (Grego & Vieira, 2005). Characterizing the spatial variability of the soil chemical properties, particularly in cultivated areas, provides relevant information for a more rational soil use and management (Montezano et al., 2006). Such studies gain particular importance in the case of mapping changes caused by different management methods (Carvalho et al., 1998; Carvalho et al., 2002). In such cases, characterizing the spatial variation of soil properties makes it possible to define differentiated management areas, which may increase the efficiency of fertilizer use and sampling schemes. Geostatistics has been used as an important and efficient strategic tool to characterize spatial variability for detailed soil analysis and description (Grego & Vieira, 2005; Silva et al., 2007). Silva et al. (2007) concluded that the amplitude of chemical soil properties in coffee plantations justifies local and differentiated fertilizer use, and that mean values do not adequately reflect nutrient demands. Marked variations in nutrient

levels and soil pH can be found frequently in cultivated fields, generally due to intensive cultivation and fertilizer use along the plantation rows (Werner, 2004).

Thus, the study hypothesis was that soil displacement due to coffee tree removal would result in spatial variations of soil chemical properties so that no spatial dependence structure could be established from the semivariograms. The purpose of this study was to characterize the spatial variability of chemical properties of a Latossolo Vermelho after bulldozing coffee trees off an over 30-year-old coffee plantation.

MATERIAL AND METHODS

This study was carried out at the Experimental Center of the Instituto Agrônomo (Santa Elisa Farm), Campinas, Sao Paulo. According to Köeppen's climatic classification, the study site is a transition between the Cwa and Cfa climatic types: a tropical dry winter, wet summer, high-altitude (± 600 m) climate. The site – a Latossolo Vermelho – had been cultivated for over 30 years with coffee. In March 1984, plants were removed by passing a bulldozer with a front blade so that yearly crops could be established. After plant removal, soil samples were taken from a georeferenced regular 20 x 20 m grid based on plot limits (Figure 1) from the layers 0–0.2 and 0.2–0.4 m; the samples were air-dried, pounded to break up clods, sieved through a 2 mm diameter mesh, and analyzed for pH, soil organic matter (SOM), exchangeable K⁺, Ca²⁺ and Mg²⁺, available P, H + Al, N-NH₄⁺ and N-NO₃⁻, as described by Raij & Quaggio (1983).

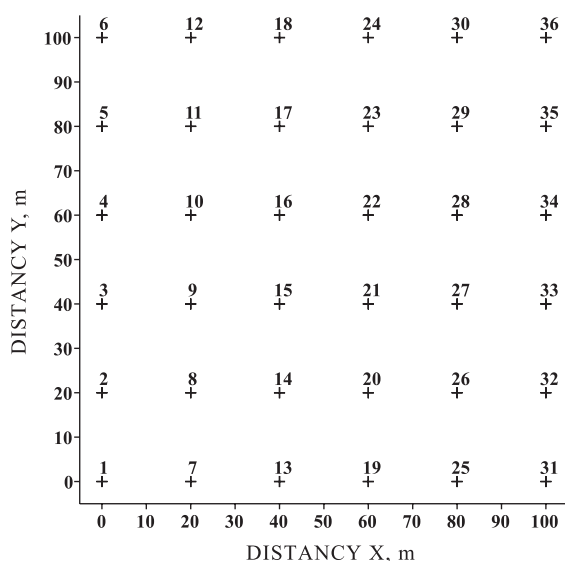


Figure 1. Sampling grid of the experimental area.

Descriptive statistical parameters (mean, variance, variation coefficients (VC), asymmetry and kurtosis) were calculated to check data dispersion and central tendency. Transformation procedures were applied to data that were initially not distributed normally. Data were analyzed by geostatistical methods (semivariogram calculation as described by Vieira, 2000) based on hypothetical intrinsic stationarity assumptions to study the spatial variability of soil properties. Semivariance $\gamma(h)$ was applied to calculate the partial autocorrelation among local neighbors, as shown in equation 1:

$$\gamma(h) = \frac{1}{2N(h)} \sum_{i=1}^N [Z(x_i) - Z(x_i + h)]^2 \quad (1)$$

where $N(h)$ is the number of pairs of measured values $Z(x_i)$ and $Z(x_i + h)$, separated by a vector h , which is the distance defined by the coordinates $Z(x_i)$ and $Z(x_i + h)$.

The spherical mathematical model was applied to spatially dependent semivariograms (equation 2).

$$\gamma(h) = C_0 + C_1 \left[\frac{3}{2} \cdot \left(\frac{h}{a} \right) - \frac{1}{2} \cdot \left(\frac{h}{a} \right)^3 \right]; 0 < h < a \quad (2)$$

$$\gamma(h) = C_0 + C_1; h > a$$

where C_0 = nugget effect, the value of $\gamma(h)$ when $h = 0$, representing variation not explained by the model or by random variation; a = reach of spatial dependence; $C_0 + C_1$ = level, the value of $\gamma(h)$ based on the reach, and C_1 = structural variance, the difference between the level and the nugget effect. Equation (2) generates $\gamma(h)$ values for the distances h ; according to Vieira (2000), measurements done in close areas are expected to be similar, compared to measurements in more distant areas. Thus, $\gamma(h)$ will increase with h until reaching a maximum value,

after which it stabilizes at a level that corresponds to the distance limit of spatial dependence, the reach. Measurements over longer distances than the reach are randomly distributed, and are thus independent from each other. The degree of spatial dependence (GD), which measures the degree of the nugget effect (C_0) relative to the level ($C_0 + C_1$) (Cambardella et al., 1994), was applied to express the spatial dependence of a variable, and was calculated using equation (3).

$$GD = \left(\frac{C_0}{C_0 + C_1} \right) \cdot 100 \quad (3)$$

According to Cambardella et al. (1994), GD can be classified into strong ($GD < 25\%$), moderate ($26\% < GD < 75\%$), and weak spatial dependence ($GD > 75\%$). After the spatial autocorrelation among samples had been demonstrated by semivariogram analysis, values with no tendencies and minimal variation were interpolated for non-sampled sites by the ordinary kriging method described by Vieira (2000). Descriptive statistics, semivariograms and kriging parameters were calculated as explained by Vieira et al. (2002). Contour maps were drawn using the SURFER 7.0 (Golden Software, 1999) software for handling and visualizing the spatial distribution of soil properties.

RESULTS AND DISCUSSION

Descriptive statistical analysis revealed that P, K, Ca, N-NH₄, and N-NO₃ contents in both layers were not normally distributed (Gaussian distribution), as shown by the Kolmogorov-Smirnov test D coefficient, at 5%; rather, asymmetry coefficient values were above 1.0 (Table 1). According to Webster (2001), asymmetry values between 0.5 and 1.0 have a lognormal frequency distribution. Because logarithmic transformation did not adequately adjust all variables (data not shown), and a normal distribution is not needed for geostatistical analysis, outlying data – over 1.5 times the interquartile interval – were excluded (Figures 2 and 3). A similar procedure was applied to data from the 20–40 cm layer (Figure 3). The approximately normal adjustment was considered enough to avoid any distribution with very long tails and eventual issues with mean kriged values.

Based on the variation coefficient (VC) classification proposed by Gomes (1984), the only low VC soil property was pH (below 10%) in both layers, supporting conclusions of Souza et al. (2004), Pontelli (2006) and Silva et al. (2007). This finding, however, was expected since this property has low variability. The VC of soil organic matter (SOM) reached around moderate values ($10\% < CV < 20\%$) in both sampled layers; the potential acidity (H + Al) in the 0–0.2 m layer had a high VC ($> 30\%$). The remaining chemical properties had very high VCs, ranging from 35% to

Table 1. Descriptive statistics of soil chemical properties

Variables	Unit	n	Mean	Var	SD	VC	C _{sk}	C _k	Prob. D _{K-S}
0.0–20 cm									
P*	mg dm ⁻³	34	15.7	65.7	8.09	51.4	0.8	0.3	> 0.15
pH	s/unit	36	4.5	0.1	0.3	6.0	1.3	1.9	> 0.15
MO	g dm ⁻³	36	47.3	59.1	7.7	16.3	1.2	2.1	0.06
K	mmol _c dm ⁻³	36	3.9	4.8	2.2	56.4	0.4	-1.2	0.10
Ca*	mmol _c dm ⁻³	34	20.8	51.0	7.14	34.3	0.9	0.1	0.02
Mg	mmol _c dm ⁻³	36	8.1	8.3	2.9	35.5	0.7	0.3	> 0.15
H + Al	mmol _c dm ⁻³	36	57.1	242.5	15.6	27.3	-0.6	-0.1	0.14
N-NH ₄	mg g ⁻¹	36	9.1	32.8	5.7	62.7	1.2	0.6	0.06
N-NO ₃ *	mg g ⁻¹	34	9.8	62.7	7.92	81.0	0.9	0.0	0.03
20–40 cm									
P*	mg dm ⁻³	33	3.8	1.2	1.11	29.2	0.4	-0.4	0.01
pH	s/unit	36	4.5	0.1	0.4	8.3	0.5	-0.6	> 0.15
MO	g dm ⁻³	36	28.1	20.2	4.5	16.0	0.1	-1.2	0.106
K*	mmol _c dm ⁻³	31	1.2	1.4	1.18	99.1	1.3	0.9	< 0.01
Ca	mmol _c dm ⁻³	36	19.7	122.8	11.1	56.4	0.7	-0.3	> 0.15
Mg	mmol _c dm ⁻³	36	5.3	3.7	1.9	36.4	0.7	1.2	> 0.15
H + Al	mmol _c dm ⁻³	36	48.6	330.8	18.2	37.4	0.4	-0.8	> 0.15
N-NH ₄ *	mg g ⁻¹	35	7.2	13.5	3.7	7.6	0.8	-0.9	< 0.01
N-NO ₃ *	mg g ⁻¹	32	2.0	2.8	1.7	85.0	1.4	1.6	< 0.01

Var: sample variance; SD: standard deviation; VC: variation coefficient; C_{sk}: asymmetry coefficient; C_k: curtosis coefficient; D_{K-S}: probability of the Kolmogorov-Smirnov test for normal distribution ($p > 0.05$).

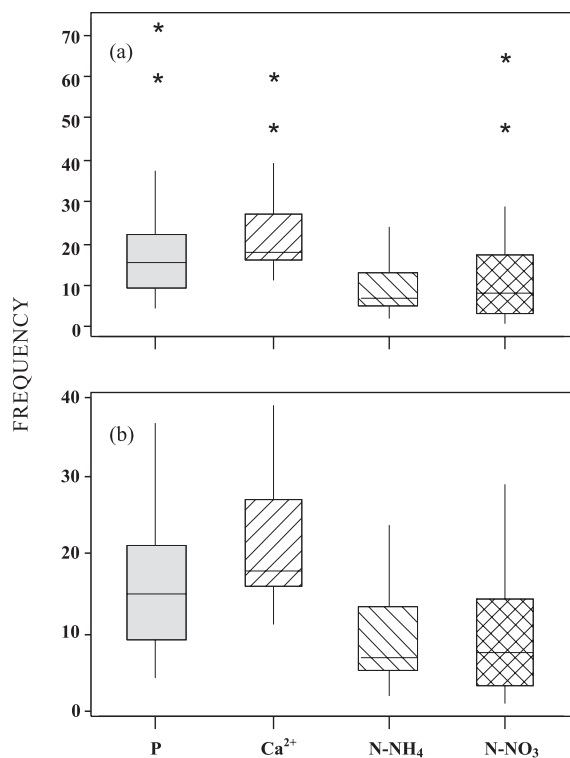


Figure 2. Boxplot of data from the 0–20 cm layer with outlying data (a) and after removal of such data (b).

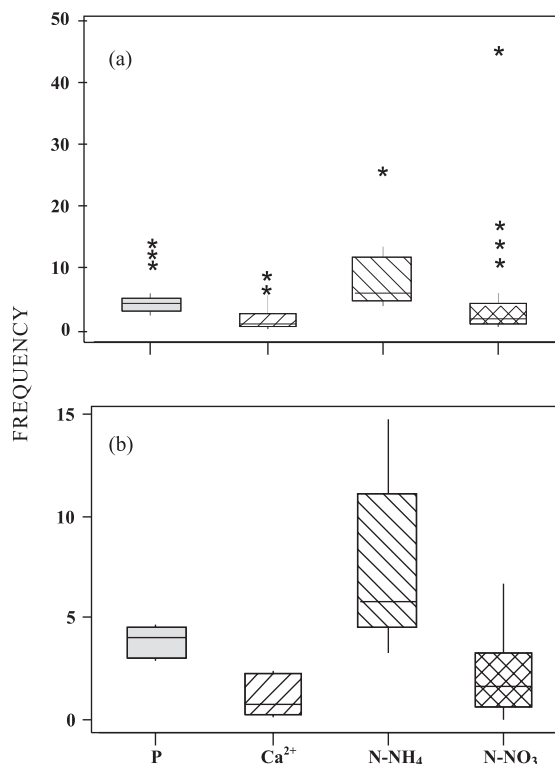


Figure 3. Boxplot of data from the 20–40 cm layer with outlying data (a) and after removal of such data (b).

about 190 %. The reach of N-NO₃ was wider than of other values in both layers: VC = 108.6 % in the 0–0.2 m layer and 189.9 % in the 0.2–0.4 m layer. Being an anion, NO₃⁻ is more mobile because of repulsion forces from predominant negative electrical charges in weathered and acid soils, typical of the soil used in this study. Many factors may cause high VC values: residual effects from previous fertilization, type of sampling scheme, and/or exposure of nutrient-poor soils due to surface erosion (Salviano et al., 1998; Montezano et al., 2006; Cavalcante et al., 2007). The expectation was that the soil movement caused by coffee tree removal would cause soil property variations at a scale that would not allow the establishment of any spatial correlation structure.

All chemical properties studied had a defined spatial dependence; semivariograms were adjusted to the spherical mathematical model (Table 2), except for pH and Mg values, which had a pure nugget effect (PNE) in the 0.2–0.4 m depth layer. The semivariograms were adjusted to the spherical model since this model suits soil property analysis best (Carvalho et al., 2003; Cavalcante et al., 2007; Zanão Junior et al., 2007; Siqueira et al., 2008); it also invalidates the hypothesis that such an adjustment would not be possible following soil movement by coffee tree removal. A mathematical model with at least 30 data pairs per point was used for the experimental semivariogram

Table 2. Determination coefficient of the spherical model adjusted to semivariograms and spatial dependence of the model

Variable	R ²	GD (%)
0.0–0.2 m		
P	0.97	42
pH	0.83	0
SOM	0.97	37
K	0.77	19
Ca	0.99	40
Mg	0.98	40
H + Al	0.93	17
N-NH ₄	0.76	16
N-NO ₃	0.85	19
0.2–0.4 m		
P	0.89	44
pH		PNE
SOM	0.97	10
K	0.71	53
Ca	0.93	43
Mg		PNE
H + Al	0.86	26
N-NH ₄	0.84	40
N-NO ₃	0.97	37

R²: adjusted determination coefficient of the model; GD: degree of spatial dependence; PNE: pure nugget effect; SOM: organic matter.

calculations, since this amount is sufficient for a robust estimate (Landim, 2003).

The nugget effect (C₀) reflects variability not explained by semivariograms for distances below the separation distance between samples (Vieira, 2000). Lower C₀ values were found for pH, K⁺, Mg²⁺ and N-NH₄ in the 0–20 cm layer, and for SOM, K, P and N-NH₄ in the 0.2–0.4 m layer (Figures 4 and 5). Potassium and N-NH₄ had low C₀ values in both layers, indicating higher continuity of spatial variability compared to the other chemical properties (Figures 4 and 5). These elements, however, are relatively mobile in soil, and the measured spatial dependence structure is valid only for a short time period after soil sampling. Soil humidity and oxidoreduction potentials affect both K dynamics and forms of soil N (Havlin et al., 1998).

Besides these variations there may be sampling errors, local variation, analytical mistakes, and other errors that make individual identification impracticable. It may therefore be easier to compare the study variables by representing random variation as a percentage. Analysis of the C₀/(C₀ + C₁) ratio makes it possible to quantify the random component (C₀) within total variance (C₀ + C₁), and is referred to as the degree of spatial dependence (GD). The spatial dependence of all chemical properties in this study was classified as *strong* or *moderate*, according to the classification of Cambardella et al. (1994). Other authors have reported similar results, which highlights the importance of knowing spatial dependence structures (Silva et al., 2003; Machado et al., 2007; Zanão Junior et al., 2007). Strong spatial dependence (GD < 25 %) of soil properties at a depth of 0–0.2 m shows that even coffee tree removal with bulldozers did not displace the soil to such a degree as to cause a completely random spatial distribution of these properties in the study area (Table 2).

As with the findings in the 0–0.2 m layer, higher C₀ values for K⁺, Ca²⁺ and Mg²⁺ in the 0.2–0.4 m layer probably result from a local application of these inputs in the shadow of the coffee tree crown for 30 years; subsequent dilution and dispersal by water in the soil profile would result in different amounts of these substances at different sampling points within the study area (Chen & Gabelman, 2000; Caires et al., 2001).

The reach of spatial dependence (α) indicates the distance from which variance becomes constant even with further increases in sampling point intervals (Journel & Huijbregts, 1991), and may be used in defining sampling procedures more effectively. Reach intervals for Ca²⁺ and K⁺ in the surface layers were 60 and 31 m each. In the 20–40 cm layer, the reach was 40 m for P and 59 m for H + Al. Thus, the 20 m distance between samples used in this study was enough to detect the spatial variability of soil properties; furthermore, sampling at a distance of up to 30 m may be used for similar variability estimates.

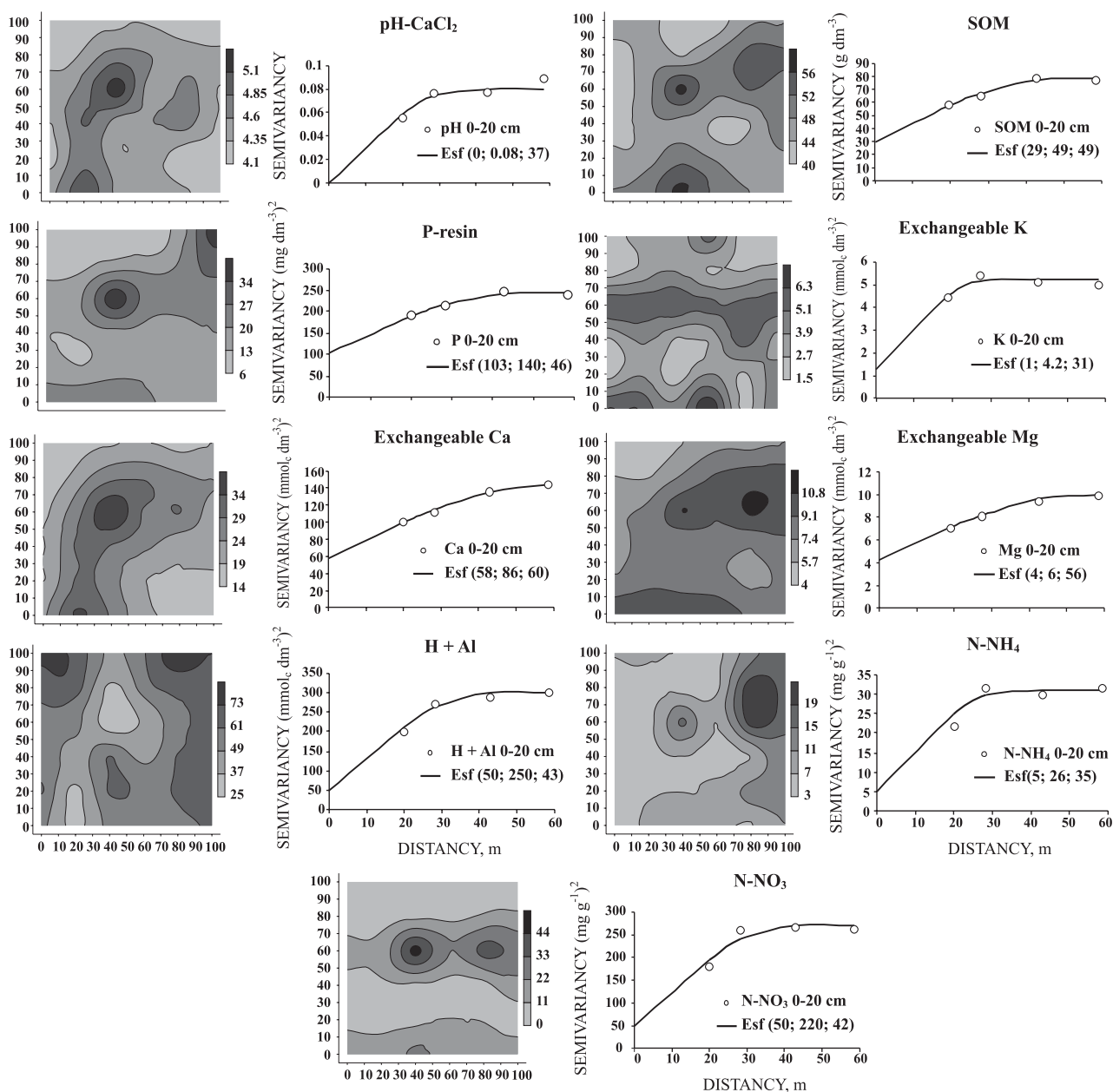


Figure 4. Spatial variability maps and semivariograms of soil chemical properties in the 0–0.2 m layer. (Modelo; C₀; C₁; a).

A visual map analysis showed that at a depth of 0–0.2 m, clear and gradual areas predominated for all soil properties (Figure 4). The pH reached from 4.1 to 5.3, with most values remaining below 4.8 in nearly the entire area, which defines soil reaction as acid. Values of P reached from 4 to 37 mg dm⁻³, showing that this nutrient was present at low and medium levels in most of the area; mean K⁺ and Mg²⁺ amplitude values reached from 1.5 to 7.5 mmol_c dm⁻³ and 4 to 12.5 mmol_c dm⁻³. The Ca²⁺ values (11 to 39 mmol_c dm⁻³) were high, according to the soil criteria for the State of São Paulo (Raij et al., 1997).

Integrated visual map analysis for both layers showed that a similarity degree between pH, SOM,

P, and Ca²⁺ maps (Figures 4 and 5). There was, however, no spatial distribution similarity between N-NH₄ and N-NO₃ in both layers. The upper right portion of the study area had higher N-NH₄ values at both depths. The N limits in the maps may vary, due to the rapid transformation of these N forms in the soil (Moreira & Siqueira, 2002).

The subsoil chemical status for root growth was diagnosed in a sample analysis of the 0.2–0.4 m layer (Raij et al., 1997). Calcium values in this layer should be above 4 mmol_c dm⁻³, whereas the Ca map analysis (Figure 5) showed that this element reached from 5 to 46 mmol_c dm⁻³, i.e., the levels of this nutrient exceeded the level considered critical for plant development by

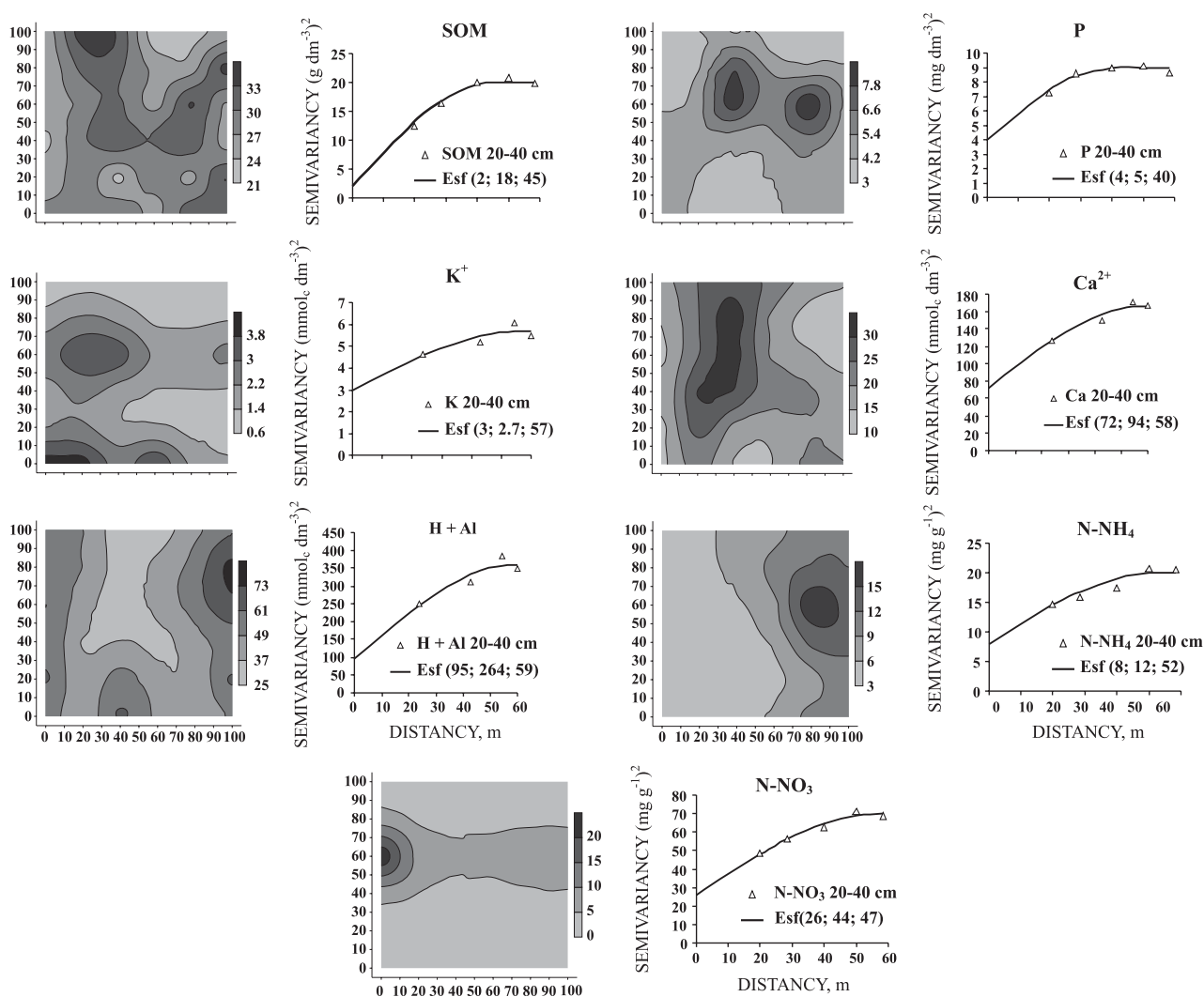


Figure 5. Spatial variability maps of soil properties in the 0.2–0.4 m layer. Model ($C_0; C_1; a$).

far. Defining areas with soil property variations of this amplitude justifies investments in soil mapping, since locally adjusted inputs may result in savings and increased productivity.

A comparison of maps for each soil property at both depths revealed that the spatial distribution of H + Al and N-NH₄ was similar. Soil displacement due to coffee tree bulldozing may have affected the spatial distribution of soil properties, as evidenced by the irregular spatial patterns of some properties; this was, however, not significant enough to affect the characterization of the spatial variability structure.

CONCLUSION

1. Removal of coffee trees with bulldozers did not result in a loss of the spatial dependence structure of the properties P, pH, SOM, K⁺, Ca²⁺, Mg²⁺, N-NH₄, and N-NO₃ in the 0–0.2 m layer.

2. The spatial dependence structure observed made it possible to identify areas with different nutrient levels by kriging in both layers.

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