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Kray, Claudio Henrique; Tedesco, Marino José; Bissani, Carlos Alberto; Gianello, Clesio; Silva, Kelly Justin da

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# TANNERY AND COAL MINING WASTE DISPOSAL ON SOIL<sup>(1)</sup>

Claudio Henrique Kray<sup>(2)</sup>, Marino José Tedesco<sup>(3)</sup>, Carlos Alberto Bissani<sup>(3)</sup>, Clesio Gianello<sup>(3)</sup> & Kelly Justin da Silva<sup>(4)</sup>

#### **SUMMARY**

Tannery residues and coal mine waste are heavily polluting sources in Brazil, mainly in the Southern States of Rio Grande do Sul and Santa Catarina. In order to study the effects of residues of chrome leather tanning (sludge and leather shavings) and coal waste on soybean and maize crops, a field experiment is in progress since 1996, at the Federal University of Rio Grande do Sul Experimental Station, county of Eldorado do Sul, Brazil. The residues were applied twice (growing seasons 1996/97 and 1999/00). The amounts of tannery residues were applied according to their neutralizing value, at rates of up to 86.8 t ha<sup>-1</sup>, supplying from 671 to 1.342 kg ha<sup>-1</sup> Cr(III); coal waste was applied at a total rate of 164 t ha<sup>-1</sup>. Crop yield and dry matter production were evaluated, as well as the nutrients (N, P, K, Ca, Mg, Cu and Zn) and Cr contents. Crop yields with tannery sludge application were similar to those obtained with N and lime supplied with mineral amendments. Plant Cr absorption did not increase significantly with the residue application. Tannery sludge can be used also to neutralize the high acidity developed in the soil by coal mine waste.

Index terms: Chromium tannery sludge, leather shavings, coal mine, heavy metal.

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<sup>(2)</sup> Professor at the Centro Federal de Educação Tecnológica, CEFET/BG. Av. Osvaldo Aranha 540, CEP 95700-000 Bento Gonçalves (RS). E-mail: ckray@terra.com.br

<sup>(3)</sup> Professor of the Department of Soil Science, Universidade Federal do Rio Grande do Sul - UFRGS. Av Bento Gonçalves, 7712, CEP 91540-000 Porto Alegre (RS).

<sup>(4)</sup> Graduate student, Escola Superior de Agricultura Luiz de Queiroz - ESALQ

# **RESUMO**: RESÍDUOS DO PROCESSAMENTO DE PELES E DE CARVÃO MINERAL APLICADOS EM SOLO

Os resíduos gerados na exploração de carvão mineral e no processamento de peles são produtos potencialmente poluentes em várias regiões do Brasil, principalmente nos Estados do Rio Grande do Sul e de Santa Catarina. O presente trabalho foi conduzido a campo com o objetivo de avaliar o efeito da aplicação e da reaplicação de resíduos carbonífero e de curtume ao Cr sobre as plantas de milho e de soja cultivadas em um Argissolo Vermelho distrófico típico, na Estação Experimental da UFRGS, localizada no município de Eldorado do Sul (RS). A primeira aplicação dos resíduos foi feita no ano agrícola de 1996/97 e a reaplicação no ano agrícola de 1999/2000. As doses de resíduos de curtume variaram conforme o pH do solo e o poder de neutralização do resíduo, tendo sido adicionadas quantidades cumulativas de Cr entre 671 a 1,342 kg ha<sup>-1</sup>. A quantidade acumulada de resíduo carbonífero foi de 164 t ha-1. Foram avaliados os efeitos das aplicações dos resíduos sobre o rendimento e a absorção de N, P, K, Ca, Mg, Cu, Zn e Cr pelas plantas. A adição de lodo de curtume propiciou rendimentos de milho e de soja semelhantes aos obtidos com a adição de N mineral e de corretivo da acidez. Não foram observadas alterações significativas na absorção de Cr pelas plantas. O efeito acidificante do solo apresentado pelo resíduo carbonífero pode ser controlado por aplicações adequadas de lodo de curtume alcalino.

Termos de indexação: cromo, lodo de curtume, metais pesados, resíduo carbonífero e serragem cromada

#### INTRODUCTION

Coal mining, hide tanning and leather product industries are important economic activities in southern Brazil. They generate, however, high amounts of residues which pose potential pollution problems. Proper handling and disposal practices are therefore required. A field experiment was started in 1996 to study the effects of such residue applications to the soil on corn and soybean crops. First results were presented by Ferreira et al. (2003). The trial was resumed with applications of the same residue types when the soil pH dropped significantly, in the 1999/2000 summer season.

# MATERIAL AND METHODS

An ongoing long-term field experiment that began in 1996 at the Federal University of the Rio Grande do Sul Experimental Station, is conducted on a Typic Paleudult soil, previously maintained under native pasture. At the beginning of the experiment, soil samples were collected at depths of 0 - 20 cm and 50 - 80 cm. The pH in the surface layer was 4.9 (1:1 water suspension); clay and organic matter contents were 250 and 25.0 g dm<sup>-3</sup>, respectively. The complete soil characterization is given by Ferreira et al. (2003).

The soil in the experimental area ( $30^{\circ} 05' 76'' S$  and  $51^{\circ} 40' 67'' W$ ) had low P and high K contents. In the 1996/97 growing season the area was prepared

for planting by plowing and disking. The residues (Table 1) were broadcast and mixed with the soil surface layer (0 to 20 cm). Tannery residues were obtained at the UTRESA (a waste-treatment station) and coal mine refuse from the COPELMI mine, near the experimental site.

The treatments (with four replications) are described in table 2. The plots (10 x 7 m) were divided to grow corn (Zea mays, L) and soybean (Glycine max, L. Merrill), sown on January 13, 2000. In the fertilized plots, 120 and 50 kg ha $^{\!-1}$  of  $P_2O_5$  (triple superphosphate) and  $K_2O$  (KCl), respectively, were added, according to regional fertilizer recommendations (SBCS/NRS, 1995). For corn, 160 kg ha $^{\!-1}$  N (urea) was also added to the treatments with N fertilization (Table 2). Soybean was properly inoculated prior to sowing. The experimental area was sprinkle-irrigated as required.

Soil samples were taken from the surface (0 - 20 cm) and subsurface (50 - 80 cm) layers 960 days after the first residue application. Similar soil samples were also taken six months after the second residue application (in July, 2000). Soil samples of the surface layer (0 - 30 cm) were taken 30 days after the second residue application from the treatments T4, T6 and T8, these samples were transported without drying to the laboratory for Cr (VI) determination (without drying), according to a method described by Bartlett & James (1996).

At the grain filling stage, four corn and 20 soybean plants of each plot were harvested to determine the dry matter production, nutrients (N, P, K, Ca, Mg, Zn and Cu) and Cr contents, according to the method described by Tedesco et al. (1995).

 $Table \ 1. \ Description \ of the \ residues \ applied$ 

Determination (1)	Tanner	y sludge	dge Leather shavings		Coal mine refuse	
Determination (7)	I(3)	$\Pi^{(3)}$	I	II	I	II
Solid content (g kg <sup>-1</sup> )	390	300	460	480	905	925
pH (water)	7.8	7.7	3.4	3.3	7.1	7.0
Organic carbon (g kg <sup>-1</sup> )	65.1	203.8	298.3	326.0	242.9	183.8
Nitrogen <sup>(2)</sup> (g kg <sup>-1</sup> )	9.8	32.5	113.1	119.7	3.6	2.3
Phosphorus <sup>(2)</sup> (g kg <sup>-1</sup> )	2.0	2.5	0.3	0.4	0.3	0.3
Potassium <sup>(2)</sup> (g kg <sup>-1</sup> )	0.10	0.12	0.10	0.12	0.07	0.06
Calcium <sup>(2)</sup> (g kg <sup>-1</sup> )	20.0	22.0	17.9	18.0	18.6	19.0
Magnesium <sup>(2)</sup> (g kg <sup>-1</sup> )	0.2	3.5	0.2	0.2	0.7	0.7
$\mathrm{Sulfur}^{(2)}$ (g kg <sup>-1</sup> )	13.0	15.0	16.5	15.9	83.6	85.0
Copper <sup>(2)</sup> (mg kg <sup>-1</sup> )	19	34	5	3	23	25
Zinc <sup>(2)</sup> (mg kg <sup>-1</sup> )	112	176	7	2	207	126
Manganese <sup>(2)</sup> (mg kg <sup>-1</sup> )	262	128	<3	<3	541	194
Sodium <sup>(2)</sup> (g kg <sup>-1</sup> )	8.4	8.1	7.0	4.3	ND	ND
Chromium <sup>(2)</sup> (g kg <sup>-1</sup> )	8.1	22.2	21.0	24.0	< 0.1	0.3
Cadmium <sup>(2)</sup> (mg kg <sup>-1</sup> )	0.14	0.14	< 0.01	< 0.01	13.02	12.85
Nickel <sup>(2)</sup> (mg kg <sup>-1</sup> )	15.0	14.5	3.1	3.2	19.8	19.2
Neutralizing value (%)	16.0	31.5	$\mathrm{ND}^{^{(4)}}$	ND	ND	ND

 $<sup>^{(1)}</sup>$ Determinations of the oven-dried material, except for pH.  $^{(2)}$ Total amount.  $^{(3)}$ Residues applied by Ferreira et al. (2003) and in this study, respectively (Kray, 2001).  $^{(4)}$ ND = not determined.

Table 2. Treatments, residue amounts applied and chromium added  $^{(1)}$ 

Treatments	First application (1996)		Second application (2000)		$\operatorname{Cr}$
	Residue	Lime	Residue	Lime	added
			t ha <sup>-1</sup> —		kg ha <sup>-1</sup>
T1 - Blank		_	_		
T2 - NPK fertilizer + lime to reach pH 6,0 (NPK+ lime)		3.4	_	6.3	_
T3 - Tannery sludge to reach pH 6 ,0 + PK (TS+PK);	21.2 TS	_	$22.4~\mathrm{TS}$	_	$671^{(2)}$
T4 - Twice the amount of tannery sludge applyed on treatment 3 + PK (2xTS+PK)	42.4 TS	_	44.8 TS	_	$1,342^{\ (2)}$
T5 - Coal waste + lime to reach pH 6.0 + NPK (CR+lime +NPK)	106.0 CR	3.4	56.0 CR	20.0	<0.02 (2)
T6 - Coal waste + tannery sludge to reach pH 6.0 + PK (CR+TS+PK)	106.0 CR 21.2 TS	_	56.0 CR 34.4 TS	_	$937^{(2)}$
T7 - Leather shavings + lime to reach pH $6.0 + NPK$ (LS+lime+NPK)	29.4 LS	3.4	30.0 LS	8.6	$1,337^{(2)}$
T8 - Cr sulfate + tannery sludge to reach pH 6.0 + PK (Cr <sub>m</sub> +TS+PK)	$21.2TS \\ 0.125~Cr_{m}{}^{(3)}$		$22.4~\mathrm{TS} \\ 0.500~\mathrm{Cr_m}$	_	1,296 (4)

 $<sup>^{(1)}</sup>$ Dry weight basis.  $^{(2)}$ Added by the residues (total amount).  $^{(3)}$ Added by  $Cr_2(SO_4)_3$ .  $^{(4)}$ Added by the residues and  $Cr_2(SO_4)_3$ .

#### RESULTS AND DISCUSSION

### Soil pH values and Cr contents

The residual effects of the first residue application were evaluated 960 days after soil sampling (Table 3). It was observed that tannery sludge was effective to maintain the pH values at a similar level to liming (T3 and T8). Higher amounts are however required to maintain an adequate pH level over longer periods (T4). The same occurs when coal mine refuse is applied, due to the acid-forming capacity of this residue by pyrite oxidation (T5, T6). Alkaline tannery sludge and acidifying coal mine refuse could therefore be applied in proper amounts to maintain adequate soil pH values for plant growth (5.5 to 6.5).

The acidifying capacity of coal residue was observed in the subsurface soil layer (50 - 80 cm) 960 days after sampling, reaching a pH value of 3.9 in treatment T5, despite lime application (Table 3). Liming should be applied more frequently in this case.

Chromium contents in the soil surface layer increased with tannery residue application and addition of Cr sulfate (Table 4). Chromium recovery was however low, reaching values from 23 to 43 % of the applied amounts. Low Cr recovery values are often reported (Castilhos et al., 2002; Ferreira et al., 2003; Domaszak, 2000). This can be due to the coarse particle segregation of the residues during sample preparation (Domaszak, 2000) or

to the low solubility of compound formation with soil minerals.

No Cr translocation to the soil subsurface layer was observed, in agreement with previous reports in the literature (Castilhos et al., 2002; Ferreira et al., 2003; Domaszak, 2000). Even with Cr sulfate application (T8), no Cr(VI) formation was detected in the soil surface layer.

## Grain yields and plant Cr contents

The plant dry matter production increased with residue application, with adequate liming and fertilization (Table 5). The dry matter production increase was however lower in the treatments with coal waste application, due to the lower pH in the subsurface soil layer (T5 and T6), mainly in corn. A smaller increase was also stated in the treatment with LS application, probably due to P immobilization by soil microorganisms, as previously observed (Ferreira, 2002). Grain yields followed the same general pattern observed for plant dry matter production.

It was observed that Cr translocation to the plant tops was low, in the mean 3.5 times lower in the grain than in the dry matter (0.30 and 1.05 mg kg<sup>-1</sup>, respectively) (Figure 1), as previously reported in the literature (Castilhos et al., 2002; Ferreira et al., 2003). The determined values were within the range usually found in plants growing on uncontaminated soil (Kabata-Pendias & Pendias, 1986; Pais & Benton-Jones 1997).

Table 3. pH values (1:1 water suspension) of the soil top layer (0-20 cm depth) at three sampling times (two after the first residue application and one after the second) and of the soil subsurface layer (50-80 cm depth) at two sampling times

	Afte	er first applica	tion	After second	After second application		
Treatment	0-20	cm	50-80 cm	0-20 cm	50-80 cm		
	180 days	960 days	960 days	180 d	ays		
1 - Blank	5.2 c A	4.5 c B	4.4 a	4.9 d AB	4.6 a		
2 - NPK + lime	5.9 ab A	5.1 ab B	4.4 a	6.2 bc A	4.6 a		
3 - TS + PK	6.0 a B	5.1 ab C	4.3 a	6.7 ab A	4.7 a		
4 - 2xTS + PK	6.2 a B	5.4 a C	4.3 a	7.1 a A	4.7 a		
5 - CR+lime + NPK	$5.0~\mathrm{c}~\mathrm{A}$	3.4 d B	3.9 b	5.3 d A	4.3 b		
6 - CR + TS + PK	6.3 a A	4.4 c B	4.1 b	6.6 ab A	4.4 b		
7 - LS + lime + NPK	$5.4 \ \mathrm{bc} \ \mathrm{AB}$	4.7  bc B	4.3 a	5.9 c A	4.6 a		
$8 - Cr_m + TS + PK$	5.9 ab B	5.1 ab C	4.4 a	6.4 bc A	4.6 a		

Means followed by the same lower case letter, in a column (treatments) and by the same upper case letter, in a row (sampling times) were not significantly different by the Tukey test (p<0.05).

Table 4. Chromium contents<sup>(1)</sup> of the surface soil layer (0-20 cm) at three sampling times (one after the first residue application and two after the second) and of the subsurface layer (50-80 cm depth) at two sampling times

	After first a	application	After second application		
Treatment	960 days		30 days	180 days	
	0-20 cm	50-80 cm	0-20 cm	0-20 cm	50-80 cm
_			mg kg <sup>-1</sup>		
1 - Blank	14 b A	14	10 d A	12 d A	13
2 - NPK + lime	12 b A	16	14 d A	15 d A	13
3 - TS + PK	68 ab B	14	114 c AB	$132~\mathrm{c~A}$	14
4 - 2xTS + PK	112 a B	16	179 b A	181 bc A	14
5 - CR+lime + NPK	12 b A	14	12 d A	11 d A	13
6 - CR + TS + PK	115 a B	16	176 b A	194 b A	13
7 - LS + lime + NPK	95 a B	17	$135~{ m bc}~{ m AB}$	151 bc A	14
$8 - Cr_m + TS + PK$	98 a B	16	290 a A	279 a A	14

<sup>(1)</sup> Extracted with concentrated HNO<sub>3</sub> + HClO<sub>4</sub> acids.

Averages followed by the same lower case letter, in a column (treatments) and by the same upper case letter, in a row (sampling times) were not significantly different by the Tukey test (p<0.05).

Table 5. Plant dry matter production and grain yield in the 1999/2000 growing season

TD	Dry ma	Dry matter		Yield		
Treatment -	Soybean	Corn	Soybean	Corn		
	$_{ m g\ plant^{-1}}$		$ m t~ha^{-1}$			
1 - Blank	6.80 b	82.9 c	1.24 b	3.27 c		
2 - NPK + lime	12.33 a	162.4 ab	2.78 a	8.60 a		
3 - TS + PK	13.78 a	176.9 a	2.81 a	8.61 a		
4 - 2 xTS + PK	13.37 a	175.4 a	2.83 a	8.71 a		
$5$ - CR+lime + NPK $^{(2)}$	10.93 a	138.6 b	2.82 a	6.91 b		
6 - CR + TS + PK	12.78 a	152.0 ab	2.87 a	8.70 a		
7 - LS + lime + NPK	11.43 a	169.1 ab	2.76 a	6.90 b		
$8 - Cr_m + TS + PK$	11.30 a	171.8 a	2.88 a	8.50 a		

Means followed by the same letter, in a column, were not significantly different by the Tukey test (p<0.05).

According to the State legislation of Rio Grande do Sul, up to 1,000kg ha<sup>-1</sup> of Cr per hectare may be applied in tannery residues (Rodrigues et al., 1993). Higher amounts were applied here (treatments T4 and T7), with no harmful effects. There is no agreement between legislations on the maximum allowable Cr addition to the soil with organic residues. While some normative institutions do not mention maximum values (USEPA, 1996; CEC,

1986), others have conservative limits, as the Brazilian CONAMA (2006), which allows a Cr application of up to  $154~\rm kg~ha^{-1}$  with sewage sludge. The naturally occurring Cr contents in several soils in Brazil exceed this limit.

In the fertilized treatments (T2 to T8), plant nutrient contents (N, P, K, Ca, Mg, Cu and Zn) in the dry matter were in the range considered adequate for plant growth (SBCS/NRS, 2004).

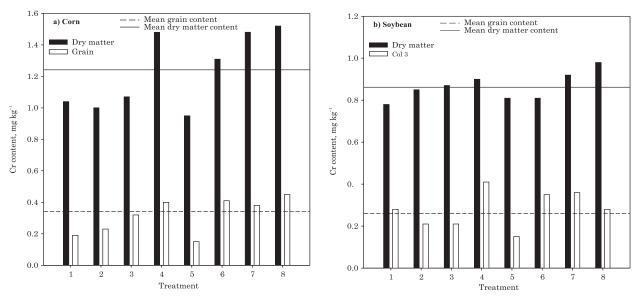


Figure 1. Plant dry matter (after flowering) and grain chromium contents.

#### CONCLUSIONS

- 1. Alkaline tannery sludge can be efficient in neutralizing soil acidity and represent a source of some plant nutrients, mainly N.
- 2. Chromium translocation to the plant tops was minor, and even smaller to grain. Cr(III) was very immobile in the soil and no oxidation to Cr(VI) was observed, even when Cr sulfate was added to the soil with tannery sludge.

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