Phosphorus Vertical Migration in Aquic Brown Soil and Light **Chernozem Under Different Phosphorous Application Rate:** A Soil Column Leaching Experiment

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Received: 21 December 2007/Accepted: 8 October 2008/Published online: 22 October 2008 © Springer Science+Business Media, LLC 2008

Abstract A soil column leaching experiment was conducted to study the vertical migration of phosphorus in aquic brown soil and light chernozem under different phosphorus fertilization rates. The results showed that total dissolved phosphorus concentration in the leachates from the two soils was nearly the same, but dissolved inorganic phosphorus concentration was obviously different. In all fertilization treatments, aquic brown soil had a higher content of phosphorus in calcium chloride extracts compared with light chernozem. But Olsen phosphorus content was higher at the soil depth beneath 0-20 cm, and increased with increasing phosphorus application rate.

Keywords Phosphorus vertical migration · Phosphorous application rate

Phosphorus (P) is an essential element for plant growth. In recent years, chemical P fertilization in China has been increasing, bringing about a potential risk of P losses from cultivated soils. Although surface runoff is considered to be the main pathway of P losses, leaching has also attracted increasing attention (Sharpley et al. 2007; Yang et al. 2007;

Presented at the 2nd International Conference on Pollution Ecology, November 4-8, 2007, Guilin, Guangxi, China.

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Brock et al. 2007). A 71 times higher P leaching loss was observed from structured clay column than from sand column after chemical P fertilization (Djodic et al. 1999), but the P leaching loss from some soils was rather low in spite of high P application rate, due to the high P sorption capacity of their subsoil (Djodjic et al. 2004). Manure application resulted in a significant increase of leachate P concentration (Kleinman et al. 2005), e.g., 0.45-0.79 mg P L⁻¹ was measured in the drainage water from manured fields (Hooda et al. 1999).

Many studies were made on the influence of manure application on soil P leaching (Chrysostome et al. 2007; Brock et al. 2007; Chardon et al. 2007), but Chinese farmers are more used to apply chemical fertilizers. Therefore, to study the vertical migration of P in our cultivated soils under chemical P fertilization is of significance in approaching the potential risk of soil P losses. In this paper, aquic brown soil and light chernozem, the main agricultural soils in north and northeast China, were chosen as test soils, and a soil column leaching experiment under different chemical P fertilization rate was conducted to monitor the contents and dynamics of different P forms in leachate, examine the downward movement of soil P, and compare the P leaching characteristics of test soils.

Materials and Methods

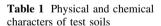
Surface soil (0-20 cm) and subsurface soil (20-60 cm) samples were collected from the aguic brown soil at Luancheng Agro-Ecological Experimental Station (37°53′ N, 114°40′ E, elevation 50 m) of Chinese Academy of Sciences in Hebei Province, and from the light chernozem in Baicheng County (45°51′ N, 123°11′ E, elevation 134 m) of Jilin Province, with their basic physical and chemical

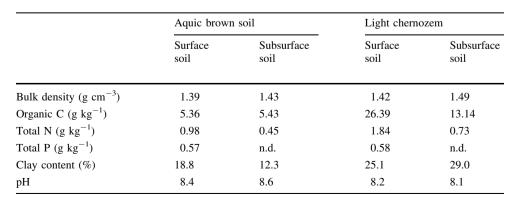


characters listed in Table 1. The samples were air-dried and 2 mm- sieved.

A set of thick-walled plexiglass columns (70 mm in inner diameter and 700 mm in length) was mounted vertically on a metal frame (Fig. 1). A hole was drilled into each column's end-cap for leachate collection, and other six holes were equidistantly set on the side-wall of each column for soil digging. A piece of nylon filter screen was set at the bottom of each column, and then, 150 g acidwashed sand was slowly packed into the column. Another piece of nylon filter screen was set above the sand, and 2 kg air-dried subsurface soil was slowly packed into and slightly pressed until its bulk density reached about 1.3 g cm⁻³. After then, 1 kg surface soil applied with trisuperphosphate (TSP) at the rates of 0, 100, 200, 400, 800, and 1,600 kg P hm⁻² (designated as treatments P0, P100, P200, P400, P800, and P1600, respectively) was packed into, and also, slightly pressed until its bulk density reached about 1.3 g cm⁻³. Each treatment was triplicated. The soil packed in column was wetted with distilled water by slow converse flow until completely water-saturated. Each day, 400 mL (10.4 cm in depth) distilled water was added to each soil column from the top, and each column was required to stop leaching by clipping the tube when there was a 5 cm water layer on the top of the column. The leaching experiment lasted for 31 days, and soil samples were dug from the holes on the side-wall at the depths of 0, 10, 20, 30, 40, 50, and 60 cm, respectively, when the leaching experiment was completed.

The leachate from light chernozem was collected every other day, and that from aquic brown soil was collected at day 1, 2, 3, 4, 8, 10, 18, and 31. After collection, the leachate was filtered with a 0.45-µm millipore membrane, and then stored at 4°C in dark for further analysis. Its total dissolved P (TDP) concentration was determined colorimetrically according to the method of Murphy and Riley (1962) after a Kjeldahl digestion (Taylor 2000), dissolved inorganic P (DIP) concentration was determined colorimetrically according to the method of isobutanol-molybdenum blue (Lu 2000), and dissolved





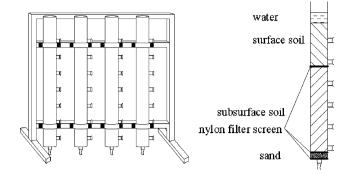


Fig. 1 Schematics of soil columns used for leaching experiment

organic P (DOP) was calculated from the difference between TDP and DIP. Soil samples dug from the columns were air-dried, ground to <2 mm, and analyzed for 0.01 M CaCl₂-extractable P (CaCl₂-P) (McDowell and Sharpley 2004) and 0.5 M NaHCO₃-extractable P (Olsen-P) (Olsen et al. 1954). All data were subjected to statistical analysis of variance (ANOVA) in the SPSS 13.0 statistical package.

Results and Discussion

Figure 2 showed that in all treatments, the TDP and DIP in the leachate from light chernozem had the same variation trend, i.e., decreased with time. The TDP and DIP were the maximum on day 1, with an average of 0.43 mg L^{-1} and 0.20 mg L^{-1} , and decreased sharply from day 1 to day 19 and from day 1 to day 15, respectively, with an average decreasing rate being 0.014 mg L^{-1} day $^{-1}$ for TDP and 0.006 mg L^{-1} day $^{-1}$ for DIP. Thereafter, the TDP and DIP decreased slowly over time, with the same decreasing rate of about 0.001 mg L^{-1} day $^{-1}$, and finally reached to about 0.13 mg L^{-1} and 0.05 mg L^{-1} , respectively. No significant difference was observed among the treatments, suggesting that chemical P fertilization rate had lesser effects on the concentrations of TDP and DIP in the leachate from light chernozem.



Fig. 2 Dynamics of TDP and DIP in the leachate from light chernozem

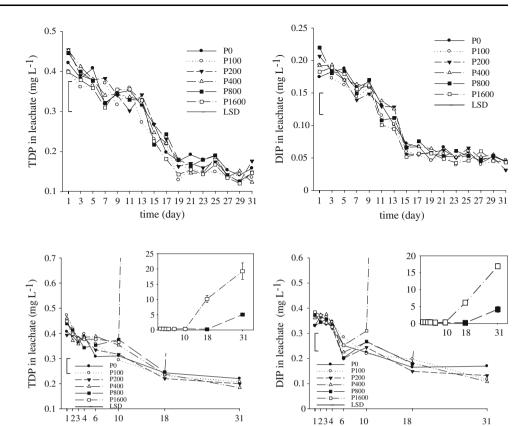


Fig. 3 Dynamics of TDP and DIP in the leachate from aquic brown soil

Somewhat different scenes were observed about the variation trend of TDP and DIP in the leachate from aquic brown soil (Fig. 3). When the P application rate was less than 400 kg hm⁻², the TDP and DIP decreased slowly with time. From the beginning to the end of the experiment, the TDP decreased from 0.43 mg L⁻¹ to 0.20 mg L⁻¹, and the DIP decreased from 0.35 mg L⁻¹ to 0.13 mg L⁻¹. In treatments P800 and P1600, an obvious increase of TDP and DIP was observed on day 31 and day 18, respectively. The concentrations of TDP and DIP in treatment P800 were 5.05 mg L⁻¹ and 4.15 mg L⁻¹ on day 31, respectively, being about 10 times of the concentrations when the experiment started, and those in treatment P1600 reached

to 20 mg L^{-1} and 17 mg L^{-1} , respectively, being about 4 times in treatment P800.

time (day)

It was found that the TDP concentration in the leachates from the two soils was nearly the same, but the DIP concentration was obviously different, being higher from aquic brown soil than from light chernozem, probably duo to the higher clay content in light chernozem. Similar result was obtained by Djodjic et al. (2004).

A comparison of Olsen-P and CaCl₂-P contents at the same depths of the two soils under different P application rate (Tables 2 and 3) showed that in light chernozem, Olsen-P content was significantly higher in upper soil layers when the P application rate was high, but had less

Table 2 Comparison of Olsen-P and CaCl₂-P contents (mg kg⁻¹) at the same depths in light chernozem under different P application rate

time (day)

	0 cm		10 cm		20 cm		30 cm		40 cm		50 cm		60 cm	
	Olsen-P	CaCl ₂ -P												
P0	21.1a	0.07a	19.8a	0.06a	13.7a	0.04a	5.9a	0.04a	4.3a	0.05a	5.0a	0.03a	4.3a	0.04ab
P100	29.2a	0.04a	30.4ab	0.05a	28.0a	0.04a	5.0a	0.03a	5.7a	0.04a	4.6a	0.05ab	5.4a	0.03b
P200	34.8ab	0.06a	47.3b	0.07a	46.1ab	0.06a	6.1a	0.05a	5.3a	0.05a	5.6a	0.04a	5.1a	0.05abc
P400	65.2bc	0.15ab	118.6c	0.25ab	91.5c	0.21b	5.9a	0.10a	4.9a	0.05a	6.2a	0.09b	5.4a	0.07bc
P800	73.2c	0.14ab	177.7d	0.36b	193.7d	0.33b	25.5a	0.20b	5.8a	0.09ab	5.2a	0.09b	6.0a	0.07abc
P1600	128.1d	0.35b	243.3e	0.65c	292.3e	0.89c	155.7b	0.87c	16.9b	0.11b	5.4a	0.09b	5.5a	0.08c

Different lowercase in a column represents significant difference at p < 0.05



0.25a

0.4a

0.15a

P200

3.7a

0.19a

13.2a

	0 cm		10 cm		20 cm		30 cm		40 cm		50 cm		60 cm	
	Olsen-P	CaCl ₂ -P												
P0	1.7a	0.06a	2.5a	0.08a	3.1a	0.25a	2.9a	0.24a	1.8a	0.18a	0.6a	0.16a	0.5a	0.14a
P100	2.8a	0.11a	7.7a	0.36a	21.2ab	0.46a	14.4ab	0.44a	2.7a	0.19a	0.9a	0.16a	0.4a	0.15a

Table 3 Comparison of Olsen-P and CaCl₂-P contents (mg kg⁻¹) at the same depths in aquic brown soil under different P application rate

0.56a 50.4c P400 8.3b 0.20a 24.6b 0.93a 1.90a 41.8c 1.79a 26.1b 1.64b 11.7b 0.67a 4.6b 0.53a 34.8b 39.0c 84.6d 4.90b 2.70b P800 18.8c 0.34a1.06a 6.34b 65.0d 4.74c 36.7c 3.89b 25.1c P1600 20.4c 0.75b67.5c 2.83b 256.2e 17.69c 94.1e 10.90c 77.2c 10.42d 70.8d 9.56c 50.2d 6.65c

29.1bc

0.77a

10.9a

0.62ab

Different lowercase in a column represents significant difference at p < 0.05

29.7b

0.93a

difference in deeper layers among all treatments. No significant difference was observed for CaCl2-P content among the treatments except P800 and P1600 that always contained more CaCl₂-P. In aquic brown soil, the contents of Olsen-P and CaCl2-P were higher in treatments P800 and P1600, but no significant differences were observed in other treatments.

Figure 4 showed that Olsen-P content was higher in light chernozem than in aquic brown soil at the depth of 0-20 cm, but mostly in adverse beneath this depth and increased with increasing P application rate. CaCl2-P content was higher in aquic brown soil than in light chernozem in the profile in all treatments. All of these suggested that aquic brown soil had a greater potential of P

1.1a

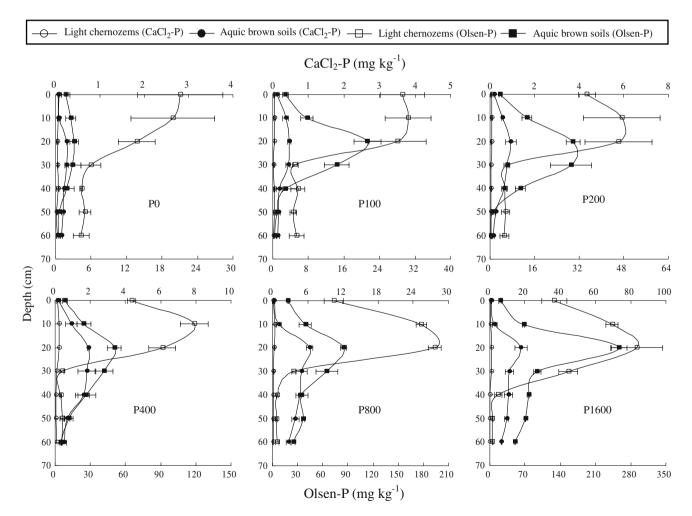


Fig. 4 Vertical distribution of Olsen-P and CaCl2-P contents in aquic brown soil and light chernozem columns under different treatments



leaching loss under chemical P fertilization, compared with light chernozem.

Acknowledgement Foundation item: The National Natural Science Foundation of China (No. 30670378).

References

- Brock EH, Ketterings QM, Kleinman PJA (2007) Phosphorus leaching through intact soil cores as influenced by type and duration of manure application. Nutr Cycl Agroescosys 77:269–281. doi:10.1007/s10705-006-9065-3
- Chardon WJ, Aalderink GH, van der Salm C (2007) Phosphorus leaching from cow manure patches on soil columns. J Environ Qual 36:17–22. doi:10.2134/jeq2006.0182
- Chrysostome M, Nair VD, Harris WG, Rhue RD (2007) Minimizing confounding factors in phosphorus leaching assessment for dairy-and poultry-manure-amended soils. Commun Soil Sci Plan 38:975–987. doi:10.1080/00103620701278054
- Djodic F, Bergstrom L, Ulen B, Shirmohammadi A (1999) Mode of transport of surface-applied phosphorus — 33 through a clay and sandy soil. J Environ Qual 28:1273–1282
- Djodjic F, Borling K, Bergstrom L (2004) Phosphorus leaching in relation to soil type and soil phosphorus content. J Environ Qual 33:678–684
- Hooda PS, Moynagh M, Svoboda IF, Edwards AC, Anderson HA, Sym G (1999) Phosphorus loss in drainflow from intensively managed grassland soils. J Environ Qual 28:1235–1242

- Kleinman PJA, Srinivasan MS, Sharpley AN, Gburek WJ (2005) Phosphorus leaching through intact soil columns before and after poultry manure application. Soil Sci 170:153–166. doi:10.1097/ 00010694-200503000-00001
- Lu RK (2000) Chemical analysis in farming soil. China Agricultural Science and Technology Press, Beijing (in Chinese)
- McDowell RW, Sharpley AN (2004) Variation of phosphorus leached from Pennsylvanian soils amended with manures, composts or inorganic fertilizer. Agr Ecosyst Environ 102:17–27. doi: 10.1016/j.agee.2003.07.002
- Murphy J, Riley JD (1962) A modified single solution for determination of phosphate in natural waters. Anal Chim Acta 27:31–36. doi:10.1016/S0003-2670(00)88444-5
- Olsen SR, Cole CV, Watanabe FS (1954) Estimation of available phosphorus in soils by extracting with sodium bicarbonate. USDA Circ. 939. US Gov Print Office, Washington DC
- Sharpley AN, Krogstad T, Kleinman PJA, Haggard B, Shigaki F, Saporito LS (2007) Managing natural processes in drainage ditches for nonpoint source phosphorus control. J Soil Water Conserv 62:197–206
- Taylor MD (2000) Determination of total phosphorus in soil using simple Kjeldahl digestion. Soil Sci Plant Anal 31:2665–2670
- Yang JY, He ZL, Yang YG, Stoffella P, Yang XE, Banks D, Mishra S (2007) Use of amendments to reduce leaching loss of phosphorus and other nutrients from a sandy soil in Florida. Environ Sci Pollut R 14:266–269. doi:10.1065/espr2007.01.378

