# PROCEEDING The 6<sup>th</sup> International Symposium for the Development of Integrated Pest Management (IPM) in Asia and Africa





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#### Proceeding

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#### PREFACE

To date, the international standing committee had arranged five international symposiums to extend IPM research successfully. In the 6th IPM symposium, we have four sessions for oral presentation and one session for poster presentation. The IPM technology contains various topics of different field so there is scarce opportunity to mutually exchange the knowledges about IPM technology in the symposiums divided into ordinary agricultural sciences. Therefore, the aim of this symposium is to exchange our knowledge of IPM, to develop the expertise for crop production, and to solve the agricultural problems using IPM.

I also sincerely appreciate all of the participants who come from the different countries far from Niigata, such as Turkey, Vietnam and Indonesia, and also appreciate the participants from Japan. Prof. Fuji comes from Akita prefectural university.

We wish to share the opportunity to recognize the current situations surrounding IPM and develop collaboration among the universities and institutes in Asia and Africa.

Dr. Keiichi Okazaki, Chairman, Local Organizing Commitee Professor, Niigata University, Japan



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## The Speech for All Participants from Dean of Academic Assembly, Institute of Science and Technology, Niigata University

It's my great pleasure to welcome you all on behalf of Niigata University, Japan to the  $6^{th}$  International Symposium for the Development of Integrated Pest Management (IPM) for sustainable agriculture in Asia and Africa 2016.

IPM has become very popular during the last two decades particularly in South and South-east Asian countries because it calls for the reduction of indiscriminate use of chemical pesticides. IPM is an environment friendly approach for pest management. It is also an important component for Good Agricultural Practices (GAP).

The conferences have sought out three main concepts i.e., i) to develop the crops resistance to various pests and diseases with the combination of conventional methods and gene technology, ii) to progress sustainable and environment friendly management of the soil fertility iii) to develop environment sound bio-pesticides and agricultural technologies for controlling various pests and diseases.

I would like to acknowledge that, previously the IPM organizing committee had arranged five international conferences to extend IPM research successfully. The 1<sup>st</sup> conference was held on in Chiang Mai University in Thailand at 2007, the 2<sup>nd</sup> in Hanoi, organized by VAST at 2008, the 3<sup>rd</sup> in Lampung University in Indonesia at 2009, the 4<sup>th</sup> in Bangladesh, organized by Bangladesh Agricultural University, Mymensingh at 2010, and the 5<sup>th</sup> in Malaysia, organized by Malaysian Agricultural Research and Development Institute (MARDI), Sabah at 2012. The IPM Standing Committee organized the past conferences with a view of the development of the sustainable agriculture in Asia and Africa. I think that, the achievements of those conferences were outstanding.

Agriculture in Asia and Africa largely depends on the use of chemical pesticides and chemical fertilizers; but the negative aspects of these have been recognized. For the establishment of the sustainable agriculture, we need to decrease the use of those chemicals as much as possible by replacing it with bio-pesticides, organic fertilizers and pest tolerant cultivars. However, still these shifts have not been succeeded yet as the many materials used in various scenes in the fields. This international conference has obligation to develop those materials based on the three projects that we have placed.

Finally, I would like to appreciate the contributions of the organizing committee lead by Niigata University for organizing such wonderful conference. I wish the success of the symposium.

Takuji Ohyama

Dean of Academic Assembly, Institute of Science and Technology, Niigata University, Japan

#### Phosphorus Adsorption Behavior as Affected by Compost, Iron ion, and Iron Concretion in a Highly Wheathered Soil

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#### Abstract

It is to be concerned that because of the high content of free iron (Fe) and (Fe) concretion, phosphorus (P) is generally tighly bound as Fe-P compounds in soil system found in the humid tropical region and, therefore, the available P for plant growth is very low. Beside as a source of complete essential nutrients for plant, source of energy and nutrients for soil organism, and for soil structure improvements, the use of organic compost would be able to inactivate the free iron and iron concretion in the soil so that the applied P from fertilizers would be available for plant. The objective of this study is to determine the effect of free Fe, iron concretion and compost materials on the P sorption behavior in a highly wheathered (Ultisols) soil. The study consists of five reatments: S (soil), SF(soil+FeCl<sub>3</sub>), SC (soil+concretion), SFO (soil+FeCl<sub>3</sub>+compost), and SCO (soil+concretion+compost). The organic compost has been developed from the mixed of fresh cow dung, coconut fine fiber waste, air dried solid waste powder of the monosodium glutamat (MSG) processing (about 9.5% of P2O5), husk, and chicken manure being composted for 30 days. The air dried sub-soil sample, iron concretion and compost had been grinded and passed through 0.5 mm sieve. The P sorption behavior of the treatments were constructed using Langmuir Plots. The results showed that the free Fe drastically increased the maximum adsorption-P and relative P-binding energy in to the soil colloid and reduce the amount of available P followed by iron concretion treatment. While the organic compost reduced the maximum adsorption-P and relative P-binding energy and the increased the available P. The highest maximum adsorption-P and relative binding energy were found in the SF (soil+FeCl<sub>3</sub> treatment), whereas the lowest value of maximum P adsorption and binding energy were in the SCO (soil+iron concretion+compost) treatment.

Keywords: P-sorption, binding energy, free iron, concretion, Langmuir plot, metal-organic complex.

#### Introduction

The intensive wheathering, leaching and decomposition processes make soils in the humid tropical region having very low organic matter content (<1%), pH (<5), base saturation (35%), and cation exchange capacity (CEC <15 cmol<sub>c</sub>kg<sup>-1</sup>) (Rachim and Suwardi, 1999). Continuously leaching of base cations and the interchange of dry and rainfall seasons couses the reduction–oxidation reaction of iron, aluminum, and manganese increased in the soils and these metals can occupy most of the cation exchange complex in high amounts that can toxic to plants (Hairiah *et al.*, 2006). These soils are generally developed from acid sandstone parent materials having low total phosphorus, acid condition, then, the dissolved iron and aluminum can form Fe-P and Al-P compounds which are unsoluble compounds resulting less available phosphorus in the soil.

The reduction–oxidation reaction of iron can form a tick layer of iron concretion with in the soil profiles, especially, in the B horizon that can be very close in the soil surface. The formation of iron concretion as the result of reduction and oxidation reactions due to the rising up and down alternately of the ground water level and create the red-yellow color of the various form of Fe concretions; the formation of Fe concretion and the high free iron in this type of soils can be toxic and limit the plant root growth (Darmawijaya, 1997).

It has been reported that the high amount of P sorption occurring in soil colloids associated with the content of extractablel Fe and Al, oxide or oxide hydrate content of Fe and Al, and clay content dealing with the type of soil colloids and clay minerals (Sanchez, 1976). The higher levels of these compounds and the higher the clay content in this type soil, the higher the phosphorus sorption capacity and, the greater the power of the P fixation (Sukmawati, 2011). The high content of free Fe and iron concretion can affect the phosphorus (P) being generally tighly bound as Fe-P compounds and, therefore, the available P for plant growth is very low in the soils. On the acid soils such as Ultisols, high content of free Fe will fix the P from soils having been fertilizered making it unavailable to the plant (Widjaja-Adhi and Sudjadi, 1987).

One way to limit the P fixed by Fe constituents in the soil system is using organic matter before P fertilizer. Beside as a source of complete essential nutrients for plant, source of energy and nutrients for soil organism, and for soil structure improvements, the use of organic compost would be able to inactivate the free iron and iron concretion in the soil so that the applied P from fertilizers would be available for plant. Organic matter has been known to suppress the solubility of Al and Fe and can decrease the amount of P adsorbed due to organic acids produced during decomposition of organic anion form that has a capacity to bind free Fe and Al in the soil solution (Schwertmann, *et al*, 1986; Stevenson and Fitch, 1986; Hakim and Sediyarsa, 1986).

The objective of this study is to determine the effect of free Fe, iron concretions, and organic material on the maximum adsorption P, the P binding energy and available P on an Ultisol soil in Lampung Indonesia.

#### Materials and Methods

Sub-soil sample used for the study was taken from the University Experiment Station of the University of Lampung, Natar. Compost had been composted of mixing of various agriculture wastes (including cow dung and chicken fresh manure, cocodust, rice husk, mud MSG waste that had been dried- powdered), FeCl<sub>3</sub> as free iron, and iron concretions taken from the Central Lampung. Soil, compost and iron concretion were grinded and sieved through 0.5 mm, Characterization of the chemical properties of soil weres pH, total-N, available-P, exchangable K, total-C, Cation Exchange Capacity (CEC), and potential-P (25% HCl).

This study consisted of five treatments: (1) soil only (S), (2) 98% of soil + 2% of FeCl<sub>3</sub> (SF), (3) 93% of the soil + 2% of FeCl<sub>3</sub> + 5% compost(SFO), (4) 80% of the soil + 20% concretions (SC), and (5) 75% of the soil + 20% of concretions + 5% of compost (SCO). Determination of P adsorbed by the colloid in the equilibrium solution was constructed using Langmuir Isotherm (Frank, *et al.*, 1998). Soil samples were each weighed 2 g and placed in a series of shake flasks, then added 20 ml of each solution series respectively. There were 6 levels of concentration of the P-solution series used: 0, 100, 250,500, 750, and 1000 mg P L<sup>-1</sup> in each having 0.01M CaCl<sub>2</sub> (the P concentration in the solution series was determined for P initial concentration before equilibration). Soil suspension shaken for 2 hours using shaker. After shaking, the suspension was centrifuged for 10 minutes at the speed of 3000 rpm, then, the clear supernatant was measured for P - equilibrium concentration by using a spectrophotometer with a wavelength of 720 nm. This procedure was conducted for five treatments, (1) pure soil, (2) soil + FeCl<sub>3</sub>, (3) soil + FeCl<sub>3</sub> + compost, (4) soil + concretion, and (5) soil + concretion + compost. Each treatment was conducted in triplicate.

Phosphorus sorption behaviour of each treatment was constructed using Langmuir Isotherm (Fox and Kamprath, 1970; Rubinger, et al., 2006) by the following equation:

C / q = (1/kb) + 1 / bC-----(1)

where: C /q = sorption index (kg L<sup>-1</sup>), k = relative P binding energy, b = maximum adsorption-P (mg kg<sup>-1</sup>), and C= equilibrium P concentration in the equilibrium solution (mg L<sup>-1</sup>).

The construction of the Langmuir Isotherm between C/q in the Y-axis and C in the X-axis to form a straight line which is equivalent to the regression equation (Y = m + nX). Values of regression equation **n** is equal to 1/b, so that the value of b can be calculated, knowing the b value, then, the value of k can also be calculated from **m** is equal to 1/kb.

#### **Results and Discussion**

#### Soil Chemical Characteristics

The chemical properties of the soil samples used in this study are presented in **Table 1**. The soil has acidic properties with a pH of 5.05. The condition of soil acidity indicates that the availability of nutrients in the soil such as phosphorus available are very low (26.6 mg kg<sup>-1</sup>). Similarly, total-N and exchangeable K are low (0.06% and  $0.24 \text{ cmol}_c \text{ kg}^{-1}$ , respectively), and total-C content is very low (0.43%). Overall the chemical properties of the soil at this particular research are low to very low. Cation exchange capacity (CEC) is also low ( $5.73 \text{ cmol}_c \text{ kg}^{-1}$ . This indicates that the soil has been intensively weathering causing low soil fertility. Having CEC value being low causing the cations in the soil such as K<sup>+</sup>, Mg<sup>2+</sup>, Ca<sup>2+</sup> and others are easily leached, consequently, plant nutrients retained in the soil are limited. Total-C being very low due to the decomposition rate is bigger the accumulation rate and stimulated by erosion and leaching. The low availability of plant nutrients, especially P, in this Ultisol Soil thought to be as affected by the high P being tightly bound to free iron and iron concretion in the soil.

#### Phosphorus and pH in the soil treated samples

Available P and pH of the soil as affect by treatments are presenting in **Table 2**. The lowest-available P content of 2.6 mg kg-1 is on the treatment of soil + FeCl<sub>3</sub> and the highest available P is the treatment of Soil + concretion+ compost (844.9 mg kg<sup>-1</sup>). This is due to the role of organic matter as a source of nutrients including P in the compost. Organic matters from compost can form organo-metal compound between organic acids with metal ions such as Fe and Al (Schwertmann, *et al*, 1986; Stevenson and Fitch, 1986) and therefore, P from fertilizers will be more available. The highest pH (6.72) was in the treatment of Soil+ concretion +compost. However, surprisingly pH of the treatment Soil + FeCl<sub>3</sub> + compost is very low (strongly acid of pH 2.97). This is due to hydrolysis of the free iron can not be neutralized by the organic material so that the pH value is very acid. Organic materials used in this study are classified alkalis pH 8.45 with available-P as much as 9.576 g kg<sup>-1</sup> and total-P 3.14% which are classified as very high (**Table 3**).

#### Maximum adsorption P and relative binding energy

To construct Langmuir Equation, the concentration of P at equilibrium conditions (C) and the value of adsorbed-P (q). The curves of the Langmuir plots of the treatments (C/ q as a function of C) are shown in **Fig. 1** and the Langmuir equation in **Table 4**. The results show that the soil treated with FeCl<sub>3</sub> and iron concretions have higher maximum adsorption P (b) and relative P binding energy P (k) compared to other treatments. The results indicate the free Fe is greatly

increasing adsorption-P in the soil. The values of maximum adsorption P in the Soil + FeCl<sub>3</sub> and Soil+concretions are 5291 and 5051 mg kg<sup>-1</sup>, respectively and its relative P binding energy are 0.0413 and 0.0300, respectively.

The results also indicate that soil treated compost can reduce the maximum adsorption-P (b) and relative P-bond energy (k). The maximum adsorption P the treatments of Soil+ FeCl<sub>3</sub> + compost and Soil + concretion + compost are 4464 and 3135 mg kg<sup>-1</sup>, respectively and the relative P binding energy are 0.0151 and 0.0094, respectively. The decreasing of P-sorption behaviour can be as a result of organic materials being able to chelate Fe (Schwertmann, *et al*, 1986; Stevenson and Fitch, 1986).

The highest maximum adsorption-P value (b) and the relative P binding energy (k) are found in the treatments of Soil + FeCl<sub>3</sub>, (b = 5291 mg kg<sup>-1</sup> and k = 0.0413), while the lowest value of maximum adsorption-P (b) and relative P binding energy (k) are in the treatment of Soil+concretion compost (3135 mg kg<sup>-1</sup> and 0.0094). While the value in the treatment Soil (only) and soil + concretions are not significantly different (4926 vs 5051 mg kg<sup>-1</sup>). Similar trend has been reported by Wijanarko and Hanudin (2010). The decreasing of the maximum adsorption-P and the relative P binding energy in the compost treatments can be as a results of organic acids (negative charge) can bind Fe (positive charge) in the soil solution before the addition of P.

#### *Significance maximum adsorption-P (b) and the relative binding energy (k)*

To determine the significance of the maximum adsorption-P (b) and the relative P binding energy, student t-test has been executed for pairs of treatments and the results are shown in **Table 5**. The results indicate that there are significantly higher, both maximum adsorption-P and the relative P binding energy when it is introduced free iron and concretion into the soils (Soil vs Soil+ FeCl<sub>3</sub> and Soil vs Soil + concretion) while the maximum adsorption-P (b) and relative P binding energy are significantly lower when it is introduced compost in the the soil systems.

#### Conclusion

The presence of the free Fe and iron concretion in the soil were significantly increasing the maximum adsorption P (**b**), while the presence of organic compost was significantly decreasing the maximum adsorption P (**b**). The presence of the free Fe in the soil was highly significant in increasing relative P-binding energy (**k**); iron concretion in soil was significantly increasing relative P-binding energy (**k**) of adsobed-P, while the presence of organic compost was highly significant in decreasing the relative P-binding energy (**k**), except the precense of free iron and compost was only significantly decreasing relative P-binding energy (**k**).

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#### Table 1.

Chemical properties of the soil Fig.1. Linear relationship between P adsorption index P (C/q) and equibrium P concentration in the solution (C)

Chemical properties	Value	Criteria*
pH H <sub>2</sub> O	5.05	Acid
Total-N (%)	0.06	Low
Available-P (mg kg <sup>-1</sup> )	26.6	Very low
Exchangeable- K ( $\text{cmol}_{c} \text{ kg}^{-1}$ )	0.24	Low
Total- C (g kg <sup>-1</sup> )	4.32	Very low
CEC ( $\text{cmol}_{c} \text{ kg}^{-1}$ )	5.73	Low
P-total– HCl 25% (g kg <sup>-1</sup> )	1.98	Low
* (Soil Research Institute, 2005).		

Table 2. Available P and pH of treated soil
---------------------------------------------

Treatment	Available P (mg kg <sup>-1</sup> )	рН
Soil	26.6 (very low)*	5.05 (acid)*
Soil+FeCl <sub>3</sub>	2.6 (very low)	2.87 (very acid)
Soil+concretion	21.8 (very low)	4.77 (acid)
Soil+FeCl <sub>3</sub> +Compost	666.9 (very high)	2.97 (very acid)
Soil+ concretion + Compost	844.9 (very high)	6.72 (Neutral)

\* (Soil Research Institute, 2005).

Sample	Available P (mg kg <sup>-1</sup> )	Total-P (%)	pH (criteria)*
Compost	9576 (very high)	3.14 (very high)	8.45 (alkalis)
Concretion	4.02 (very low)	0.004 (very low)	4.43 (very acid
1 (G 11 D	1		

Table 3. Available P ar	d pH compost	and concretion
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\* (Soil Research Institute, 2005).

Table 4.	Langmuir	linear o	equation	of the H	sorption
	Langmun	moury	equation	or the r	sorption

Perlakuan	Langmuir linear equation	$b (mg kg^{-1})$	k	r <sup>2</sup>	
Soil	C/q = 0,00901 + 0,000203 C	4926	0,0225	0,963	
Soil + FeCl <sub>3</sub>	C/q = 0,00458 + 0,000189 C	5291	0,0413	0,942	
Soil+Concretion	C/q = 0,00660 + 0,000198 C	5051	0,0300	0,937	
Soil+ FeCl <sub>3</sub> +compost	C/q = 0,01483 + 0,000244 C	4464	0,0151	0,975	
Soil+concretion+compost	C/q = 0,03412 + 0,000319 C	3135	0,0094	0,971	
Where $b = maximum$ adsorption of $\mathbf{P} \cdot \mathbf{k} = binding$ energy $\cdot \mathbf{r}^2 = box fision$ $C = solution \mathbf{P}$ and $c = c$					

Where: b = maximum adsorption of P ; k = binding energy ; r<sup>2</sup> = koefisien, C= solution P and q= adsorbed P

Table 5. Student-t test significances of maximu adsorbed P (b) and binding energy (k)

Treatment	Calculated -t		
reatment	b	K	
Soil vs Soil + FeCl <sub>3</sub>	25.21*	142.38**	
Soil vs Soil+Concretion	8.81*	12.45*	
Soil vs Soil+ FeCl <sub>3</sub> +compost	13.41*	26.57*	
Soil vs Soil+concretion+compost	19.25*	45.02**	
Soil + FeCl <sub>3</sub> vs Soil+Concretion	8.99*	15.04*	
Soil + FeCl <sub>3</sub> vs Soil+ FeCl <sub>3</sub> +compost	19.90*	59.68**	
Soil + FeCl <sub>3</sub> vs Soil+concretion+compost	27.07*	75.34**	
Soil+Concretion vs Soil+ FeCl <sub>3</sub> +compost	20.84*	58.32**	
Soil+Concretion vs Soil+concretion+compost	18.00*	65.13**	
Soil+ FeCl <sub>3</sub> +compost vs Soil+concretion+compost	15.69*	115.68**	
Where: $b = Ads$ Maximum of P; $k = binding energy ** = ve$	ry significant (1%)	; *= significant	

(5%)