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Effect of Drying on Phosphorus Distribution in Poultry Manure

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Abstract: Laboratory drying may alter manure phosphorus (P) distribution. The effects of freeze, air (22°C), and oven (65°C) drying on sequentially fractioned poultry manure P were examined. Higher drying temperatures resulted in lower percentage of dry matter. Increased H₂O- and decreased sodium bicarbonate (NaHCO₃)-extractable P with drying provided evidence that drying increases poultry manure P solubility. Labile fractions were predominantly inorganic P (P_i), whereas sodium hydroxide (NaOH) and hydrochloric acid (HCl) fractions had significant amounts of organic P (P_o). Drying altered H₂O- and NaHCO₃-extractable P_i but had no consistent effect on P_o in these fractions. This work suggests that variations due to drying should be

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taken into consideration when evaluating manures for P availability or when comparing data in which different drying methods have been utilized.

Keywords: Inorganic phosphorus, manure drying, organic phosphorus, phosphorus, poultry manure, sequential phosphorus fractionation, total phosphorus

INTRODUCTION

Animal manures are rich in plant nutrients and have traditionally served as a valuable fertilizer resource for cropland. However, modern intensive animal and crop production systems are becoming more spatially separate, leading to concentration of surplus nutrients in areas surrounding livestock operations. Effective management practices are needed to optimize manure recycling while limiting risk of adverse environmental impact from nutrient runoff. A more thorough understanding of phosphorus (P) characteristics from manure fertilizers is essential to reach this goal.

The nonuniform composition and high moisture content of most manure samples necessitate sample pretreatments (i.e., homogenization, drying, and grinding) prior to laboratory analysis. Frequently used drying processes include air drying (22°C) (Elias-Azar, Laag, and Pratt 1980; He et al. 2006a), oven drying (65 or 105°C) (Akinremi et al. 2003; Dou et al. 2000; Sharpley and Moyer 2000), and freeze drying (He et al. 2004b; Leinweber, Haumaier, and Zech 1997). Comparative investigations have shown that manure P is affected by drying process (Ajiboye, Akinremi, and Raczy 2004; Akinremi et al. 2003; Gerritse and Eksteen 1978; Sistani et al. 2001). Sistani et al. (2001), in a study on the effects of drying method on nutrient composition of manure from broiler chickens, noted that freeze or oven drying (105°C) resulted in lower inorganic P (P_i) and total P (P_{tot}) concentrations than air drying. Akinremi et al. (2003) examined P_{tot} concentrations of freeze-dried, air-dried (30°C), and 105°C-dried organic amendments (poultry, cattle, and pig manure) and found that 105°C-dried samples had lower P_{tot} concentrations than manures dried under other conditions. In addition, Chapuis-Lardy et al. (2004) reported lower water-soluble P_i concentrations in dried dairy manure than in fresh wet feces from a diverse group of commercial dairy farms in the northeastern United States.

Phosphorus (P_i) solubility is primarily related to the solubility constants of mineral phosphates, and temperature may influence particular products formed (Gerritse and Eksteen 1978). Effects of drying on organic phosphorus (P_o) are not as clear. Solubility of P_o in pig slurry was not affected by drying (0–100°C) and redispersion in water (Gerritse and Eksteen 1978). Furthermore, freeze- and 65°C-dried samples of broiler manure had only a slight decrease in P_o , whereas P_i content decreased up to 37% with drying (Sistani et al. 2001).

Sequential fractionation has been used to investigate P forms in soils and manures, as well as their transformation under various environmental

conditions (Ajiboye, Akinremi, and Racz 2004; Akinremi et al. 2003; He, Griffin, and Honeycutt 2004a, 2004b; He et al. 2006a; Hedley, Stewart, and Chauhan 1982; Leinweber, Haumaier, and Zech 1997; Schlichting and Leinweber 2002). This method applies a series of extractants, such as water (H_2O), sodium bicarbonate ($NaHCO_3$), sodium hydroxide ($NaOH$), and hydrochloric acid (HCl), to functionally separate P into fractions according to lability. Thus, sequential fractionation enables characterization of multiple P forms, providing greater insight into how drying processes affect P distribution than would P determination with a single extraction or digestion. Using this procedure, Ajiboye, Akinremi, and Racz (2004) characterized the influence of $105^\circ C$ drying on P forms in different animal manures. Drying at $105^\circ C$ increased H_2O -extractable P_i in both swine and dairy manures. In swine manure, the major contributor to the increase was hydrolysis of H_2O -extractable P_o to H_2O -extractable P_i ; however, in dairy manure, the transformation occurred between fractions, with both $NaHCO_3$ -extractable P_i and $NaHCO_3$ -extractable P_o converted to H_2O soluble forms. In addition, changes have been found in the more stable P forms of manure with drying. Schlichting and Leinweber (2002) investigated the effects of drying on sequentially extracted P fractions from organic peat soils. Drying led to substantial increases in residual P concurrent with decreases in $NaHCO_3$ - and $NaOH$ -extractable P_o fractions.

Poultry manure is an excellent source of plant nutrients, including P. Evaluation of the effect of drying on poultry manure P distribution will improve accuracy in characterization of this type of manure, a critical step in developing sound P-management practices. This study was conducted to compare P characteristics of poultry manures after freeze drying, air drying ($22^\circ C$), and oven drying ($65^\circ C$). In addition, to find an efficient method for evaluating plant-available P in poultry manure, the relationship of P extracted by a single sodium acetate buffer (100 mM, pH 5.0) with P in sequentially extracted fractions was evaluated.

MATERIALS AND METHODS

Manures

Fresh poultry manure samples were collected from caged layer hens at two commercial egg farms in central Maine. A total of three manure samples were collected; PM-1, from layer house with daily waste removal; PM-2, layer house with a single annual cleaning; and PM-3, house with 13-week-old growing birds that received cleaning weekly. In addition to fecal material, the manure waste contained some feathers and dropped feed. For each of the three manures, five grab samples of manure were removed from inside of the poultry house, pooled, and stored in an airtight container at $4^\circ C$. Manures were then homogenized using a food processor. Undried

Table 1. Characteristics of poultry manures used in this study

Manure	P	Fe	Al	Mn	Ca	Mg	C	K	EC (ds m ⁻¹)	pH
	g kg ⁻¹ dry matter ^a									
PM-1	13.9 ± 0.6 ^b	1.14 ± 0.11	0.53 ± 0.02	0.27 ± 0.01	153 ± 5	6.89 ± 0.12	335 ± 6	26.0 ± 1.7	15.1	6.82
PM-2	19.0 ± 2.1	2.54 ± 0.36	1.04 ± 0.17	0.36 ± 0.04	152 ± 11	11.0 ± 0.4	278 ± 14	34.6 ± 3.7	12.5	7.96
PM-3	27.8 ± 3.5	1.65 ± 0.12	0.78 ± 0.08	0.54 ± 0.07	70.5 ± 1.4	11.3 ± 1.5	379 ± 2	44.4 ± 0.4	20.2	7.00

^aDry matter based 105°C-dried sample data from the maine agricultural and forestry experiment station analytical laboratory.

^bAverage from three replicate samples ± standard deviation.

manures were maintained at -20°C . Selected manure properties (Table 1) were determined by the Maine Agricultural and Forestry Experiment Station (MAFES) Analytical Laboratory. Manure was dry ashed at 550°C for 6 h, and the ash was dissolved in 50% HCl. The resulting solution was analyzed by inductively coupled plasma to atomic emission spectroscopy (ICP-AES) for P, potassium (K), calcium (Ca), magnesium (Mg), iron (Fe), aluminum (Al), and manganese (Mn) using a ThermoJarrell Ash IRIS 1000. Carbon content of the manure samples was determined by combustion using a Leco CN2000 analyzer. Electrical conductivity (EC) and pH were determined using a 6:1 water (mL) to freeze-dry manure (g) ratio. A portion of each manure was desiccated via 1) oven drying at 65°C , 2) air drying at 22°C , or 3) freeze drying, until no further decrease in weight was observed. After drying, the manures were ground in a Wiley mill, sieved through a 2-mm screen, and stored at -20°C until analysis.

Sequential Fractionation

A modified Hedley sequential fractionation procedure as described by He, Griffin, and Honeycutt (2004a) was used in this study. Each sample (0.25 g, 65°C dry equivalent) was sequentially extracted in 25 mL of distilled water, 0.5 M NaHCO_3 (pH 8.5), 0.1 M NaOH, and 1 M HCl for 2 h (water) or 16 h (all others) at 22°C on an orbital shaker at 250 rpm. After each extraction, the tubes were centrifuged at $23,700 \times g$ for 30 min at 4°C . The supernatant was passed through a $0.45\text{-}\mu\text{m}$ filter (Fisherbrand MCE membrane; Fisher Scientific, Pittsburgh, PA). The NaHCO_3 , NaOH, and HCl fractions were diluted and adjusted to pH 5.0. Ethylenediamine tetraacetic acid (EDTA) (1 mM final concentration) was added to the NaOH fraction to prevent P compounds from precipitating during pH adjustment. Manure samples (0.25 g, 65°C dry equivalent) were also separately extracted with 25 mL of 100 mM Na acetate buffer (pH 5.0) for 1 h, followed by an additional extraction with 10 mL of buffer for 1 h. Centrifugation and filtration were performed as previously described. Both acetate buffer extracts were combined. All extractions were performed with nine replications.

Phosphorus Determination

P_1 was determined using a modified molybdate blue method (He and Honeycutt 2005). P_{tot} concentrations of manures and extracts were determined by ICP-AES after dry combustion at 550°C and digestion in 0.5 M sulfuric acid (H_2SO_4). Phosphorus (P_o) was estimated as the difference between P_{tot} and P_1 . Residual P was calculated as the difference between manure P_{tot} determined as described previously and the sum of P_{tot} extracted by all four extractants.

Statistical Analysis

Statistical analyses were performed using SAS (1999). One-way analysis of variance (ANOVA) was computed to examine the effects of drying and extractant. Significance of differences among treatments was determined by mean separation using Fisher's least significant difference test.

RESULTS AND DISCUSSION

Effect of Drying Processes on Dry-Matter Content

The three poultry manures contained about 24–38% dry matter (DM) when dried at 105°C (Table 2). DM content was altered by the drying procedures. The general trend was that higher drying temperatures (from 22 to 105°C) resulted in lower DM. This observation seems reasonable because temperature increases would cause interlayered water molecules to become more active and thus easier to drive off from the manure matrix. The near vacuum condition (<500 mT) created by freeze drying was as effective as high temperatures in removing manure moisture from PM-2 and PM-3 but not from PM-1. Despite the trend toward increased moisture removal with higher temperatures, overall change in DM with drying method was not consistent across manures. Drying process had little effect on DM content of PM-1, but 105°C-dried PM-2 had only 38% DM compared to 49% DM for 22°C-dried PM-2. Manure properties, such as water-holding capacity and concentration of intracrystalline water, may have contributed to this variation. As an example, exceptionally low vapor pressure maintained over samples during extensive freeze drying has the capacity to evaporate intergranular pore water but cannot release that portion of moisture present as intracrystalline water. In contrast, high temperatures associated with oven drying could decompose crystalline water and other labile compounds, leading to additional moisture loss. Scholl et al. (1997) stated that drying at 105°C more aggressively (i.e., completely) evaporated interlayered water in sediment samples than other drying

Table 2. Effects of drying on dry matter content (%)

Treatment	PM-1	PM-2	PM-3
Freeze dry	33.2a	40.3c	26.9b
Air (22°C)	33.0a	48.8a	30.5a
Oven (65°C)	30.0b	41.6b	24.7b
Oven (105°C)	30.0b	38.3d	24.0b

Note. Averages followed by the same letter for each manure are not significantly different ($\alpha = 0.05$).

methods. They also determined that oven-dried porosity values of sediment samples exceeded those of freeze-dried samples, whereas no difference was recorded between oven- and freeze-dried samples of artificial sediments (glass beads and seawater) that lack a clay component. Further validating our observation of drying effects, Akinremi et al. (2003) noted that 105°C-dried samples contained less moisture than manures subjected to other drying methods, but no difference was distinguished between freeze- and air- (30°C) dried poultry manure samples. The inconsistent impact of drying on different samples suggests caution be taken when comparing published P data in which DM is calculated based on different drying processes.

Effect of Drying on P Distribution in Sequentially Extracted Fractions

Of the P_{tot} in the undried manure samples, 11–20% was H_2O extractable, 19–37% was $NaHCO_3$ extractable, 3–8% was $NaOH$ extractable, 31–45% was HCl extractable, and 0–25% was residual (Figure 1). Water-extractable P in manure is correlated with P in runoff from recently amended soils and is increasingly used as an indicator of environmental risk (Kleinman et al. 2002, 2005). All drying methods increased the quantity of H_2O -extractable P in PM-1 and PM-3 with the greatest change in 65°C drying, followed by 22°C drying, and then freeze drying. Drying at 65°C also increased H_2O -extractable P in PM-2. Increased drying temperature could cause sorbed P to become more H_2O soluble. Previous studies have yielded conflicting results on the effects of drying on H_2O -extractable P in manures, and it appears that P solubility is related to specific manure chemico-physical characteristics (Ajiboye, Akinremi, and Racz 2004; Kleinman et al. 2002, 2005; Sistani et al. 2001). Ajiboye, Akinremi, and Racz (2004) reported 105°C-dried dairy manure contains 25% more H_2O -extractable P than undried samples but did not see similar effects in swine or beef cattle manure. In contrast, poultry manure has been found to have decreased H_2O -extractable P with air, 65°C, and freeze drying but little change with drying at 105°C (Sistani et al. 2001). Incorporation of P into aggregated particles during dehydration has been suggested as a possible cause for decreased bioavailable P in air-dried sediment samples (Twinch 1987). These differential effects of drying on H_2O -extractable P reflect the complex nature of animal manure and our associated lack of understanding this complexity.

Drying decreased P extracted by $NaHCO_3$ (Figure 1). Manures dried at 65°C and freeze dried had 20–51% and 11–47% less $NaHCO_3$ -extractable P compared to undried manure, respectively. Air drying had a similar effect with 18–28% lower $NaHCO_3$ -extractable P than undried manure. The smallest extracted fraction was $NaOH$ -P, comprising an average of 6% of total P for the manures. Small but significant decreases in $NaOH$ -extractable P occurred in PM-1 with all drying types. Differences in 65°C- and freeze-

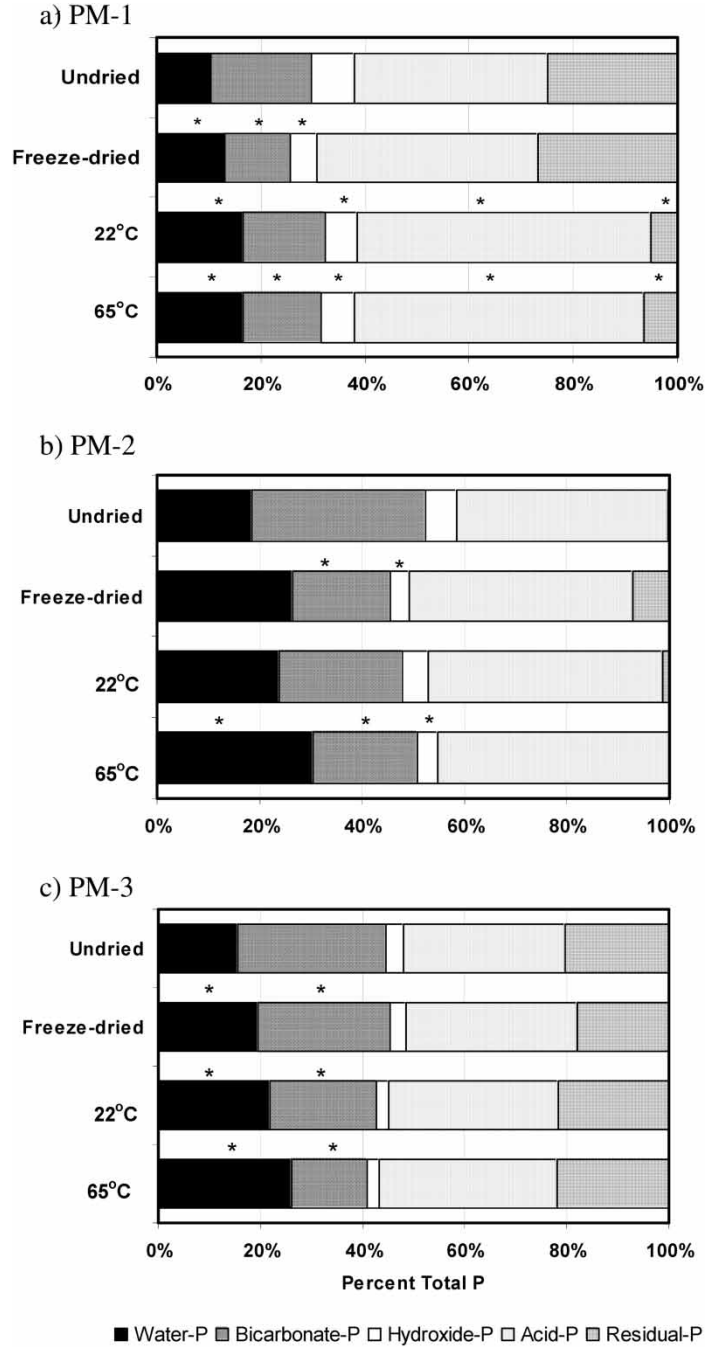


Figure 1. Effects of drying on total P distribution in sequentially extracted fractions of poultry manure. Symbol * indicates significant difference ($\alpha = 0.05$) from undried manure.

dried samples were seen in PM-2. No significant drying effect was observed on NaOH-extractable P in PM-3. Different types of organic amendments may also differ in response of NaOH-extractable P to drying. As an example, 105°C drying decreased NaOH-extractable P in beef manures and biosolids but increased it in swine manure (Ajiboye, Akinremi, and Racz 2004).

Both 65°C and 22°C drying increased HCl-extractable P in PM-1 by 33% but had no effect on PM-2 or PM-3. Residual P, the portion of P left in the manure after all extractions, was disparately affected by drying. One manure, PM-1, had about 75% less residual P after both 65°C and 22°C drying but was relatively unchanged by freeze drying. Drying had no effect on the calculated concentration of residual P in either PM-2 or PM-3. The variation associated with drying effects on less labile and residual P forms is consistent with work by Ajiboye, Akinremi, and Racz (2004), in which 105°C drying increased HCl-extractable P and decreased residual P in dairy and beef manures but had no effect on swine manure.

Transformation of P from one fraction to another as a result of sample drying has been previously reported (Ajiboye, Akinremi, and Racz 2004; Sistani et al. 2001). Heat associated with oven drying may change the solubility of some P forms, leading to elevated levels of H₂O-soluble phosphorus. Ajiboye, Akinremi, and Racz (2004) noted that oven-dried manures from dairy and swine manures have lower NaHCO₃-extractable P content than undried samples, suggesting transformation of P forms in these organic amendments. The boiling process involved with high-temperature oven drying could cause the hydrolysis of NaHCO₃-extractable P_o as well as the desorption of P that is held on the surfaces of organic and inorganic colloids in dairy manure. Desorbed P could then appear in the H₂O-extractable fraction of the oven-dried samples. Other researchers have noted a shift toward more stable and recalcitrant P forms after manure drying. McDowell and Stewart (2005) examined fresh and air-dried dung samples from grazing ruminants (dairy cattle, deer, and sheep) using sequential fractionation and solid-state ³¹P nuclear magnetic resonance (NMR) spectra, revealing that the labile P in fresh manures shifted into the more recalcitrant HCl and residual fractions with air drying.

Effects of Drying on Organic and Inorganic Phosphorus Fractions

To further evaluate the effects of drying on P distribution, sequentially extracted P was separated into P_i and P_o (Table 3). Water extracts of undried manures contained primarily P_i, comprising 73–94% H₂O- and 64–87% NaHCO₃-P_{tot}. In contrast, Sistani et al. (2001) reported that 72–83% of H₂O-extractable poultry manure P was in organic forms.

In PM-3, P_i content determined by a modified molybdenum blue method was greater than total P determined by ICP-AES for both the H₂O and

Table 3. Effects of drying on inorganic (P_i) and organic phosphorus (P_o) distribution in sequentially extracted fractions of poultry manure (g P kg⁻¹ dry matter)^a

Treatment	H ₂ O		NaHCO ₃		NaOH		HCl	
	P _i	P _o	P _i	P _o	P _i	P _o	P _i	P _o
PM-1								
Undried	1.08 ^b	0.39 ^c	1.71 ^a	0.95 ^a	0.31 ^a	0.85 ^a	3.26 ^b	1.93 ^b
Freeze dry	1.27 ^b	0.58 ^{bc}	1.09 ^c	0.63 ^a	0.24 ^a	0.46 ^b	2.67 ^c	3.29 ^a
22°C	1.50 ^a	0.83 ^a	1.52 ^{ab}	0.66 ^a	0.27 ^a	0.60 ^b	3.60 ^a	4.26 ^a
65°C	1.62 ^a	0.68 ^{ab}	1.30 ^{bc}	0.81 ^a	0.27 ^a	0.59 ^b	3.59 ^a	4.18 ^a
PM-2								
Undried	3.05 ^b	0.76 ^a	6.06 ^a	0.89 ^a	0.44 ^a	0.82 ^a	6.57 ^a	1.92 ^a
Freeze dry	4.17 ^{ab}	1.00 ^a	3.11 ^c	0.56 ^a	0.34 ^c	0.38 ^c	6.25 ^a	2.23 ^a
22°C	4.12 ^{ab}	0.90 ^a	4.55 ^b	0.51 ^a	0.40 ^{ab}	0.66 ^{ab}	7.06 ^a	2.57 ^a
65°C	5.10 ^a	1.10 ^a	3.41 ^c	0.73 ^a	0.36 ^{bc}	0.47 ^{bc}	6.69 ^a	2.52 ^a
PM-3								
Undried	4.08 ^d	0.26	8.60 ^a	—	0.71 ^a	0.23 ^a	6.68 ^a	2.48 ^a
Freeze dry	5.49 ^c	— ^c	7.23 ^b	—	0.65 ^a	0.25 ^a	6.73 ^a	2.60 ^a
22°C	6.92 ^b	—	6.17 ^c	—	0.64 ^a	0.06 ^a	6.74 ^a	2.47 ^a
65°C	7.89 ^a	—	4.58 ^d	—	0.54 ^a	0.13 ^a	7.42 ^a	2.26 ^a

^aDry matter based on 65°C-dry sample.

^bAverages followed by the same letter within columns for each manure are not significantly different ($\alpha = 0.05$).

^cCalculated results had negative values.

NaHCO₃ fractions. However, hydrolysis by phosphatase enzymes did indicate that PM-3 contained about 0.1 g P kg⁻¹ as P_o (data not shown). Disagreement between ICP and colorimetric methods of phosphorus quantification has been noted previously (Wolf 2005). The commonly used Murphy and Riley colorimetric procedure measures primarily orthophosphate P, although some organic P compounds can be hydrolyzed by acidic reagents in the assay (Dick and Tabatabai 1977; Murphy and Riley 1962). A modification proposed by Dick and Tabatabai (1977) was utilized, involving the addition of a citrate–arsenite reagent to complex with excess molybdate ions and prevent further formation of blue color from acid-hydrolyzed P in the assay. Wolf et al. (2005) proposed that higher P measurements of manure extracts determined with the Murphy and Riley colorimetric procedure were the result of color interference, as extracts can be brown or amber after centrifugation and filtration. Results from ICP and the Murphy and Riley procedure are generally correlated within 5 to 10% of each other (Wolf et al. 2005). In the current study, NaHCO₃ and NaOH extracts of our poultry manures did exhibit brown coloration. The proportion of total P in organic forms was greater in the NaOH and HCl fractions than the more labile fractions of our manures. Undried PM-1 and PM-2 had P_o concentrations representing 65% and 73% of NaOH-extractable P, respectively. In HCl fractions, 23–37% of extractable P was P_o. This observation supported the previous argument that it should not be simply assumed that sequentially extracted 1 M HCl fractions contained no organic P (He et al. 2006b).

Drying produced changes in manure H₂O-extractable P_i content. Drying at 65°C increased H₂O-extractable P_i in all three manures by 47–92% (Table 3). Freeze- and 22°C-dried samples of PM-1 and PM-3 also had higher H₂O-extractable P_i than undried manures. Drying and grinding have been reported to change P_i solubility in dairy manure, resulting in lower H₂O-extractable P_i in 65°C-dried material (Chapuis-Lardy et al. 2004). Very little change was observed in H₂O-soluble P_o with drying. In one manure, PM-1, there was a slight increase in P_o with 22°C and 65°C drying. This is consistent with other reports of H₂O-extractable P_o being relatively unaffected by drying (Akinremi et al. 2003).

Drying also had an effect on NaHCO₃-extractable P_i, with as much as an 80% decrease in 65°C-dried manure. Magnitude of the drying effect was different between manures, but 65°C and freeze drying had significant effects on NaHCO₃-extractable P_i from all three manures. There was no effect of drying on NaHCO₃-extractable P_o. NaOH-extractable P_i was largely unaffected by drying, with the exception of a reduction from 0.44 to about 0.35 g P_i kg⁻¹ in PM-2 due to 65°C drying. Organic P extracted by NaOH was reduced by all drying types in PM-1 (up to 85%) and by freeze and 65°C drying in PM-2. McDowell and Stewart (2005) found that NaOH-extractable P_o was the only fraction of P_o to significantly change in air-dried compared to undried animal dung. In this study, HCl-extractable P_i in PM-1 was reduced by freeze drying but slightly increased by 22°C and

65°C drying. In addition, all drying methods resulted in higher HCl-extractable P_o concentrations in PM-1 compared to undried manures. No effect was observed of drying on HCl-extractable P in either PM-2 or PM-3.

Analysis using ^{31}P NMR on animal dung by McDowell and Stewart (2005) showed that air drying changes P_o concentration, possibly due to degradation of diesters. Gerritse and Eksteen (1978), in a study using high performance liquid chromatography (HPLC) analysis of gel filtrations of swine manure, observed that desiccation by oven (60 and 100°C) and freeze drying did not result in an overall change in the quantity of manure P_o present. However, these researchers did find that high-molecular-weight P_o decreased after oven drying and subsequent rewetting. Increases in P_i solubility as well as transformation of P_o from the NaOH to the HCl fraction in our poultry manures further the view that manure P cannot be assumed to behave equivalently after drying. Summation of P_i and P_o values from the sequential fractions in Table 3 revealed that total P concentration of PM-1 and PM-2 did not change significantly with drying (data not shown). It is assumed that incorporation of P_o into stable aggregates during the drying process is the cause of increased P_o stability rather than heat-associated degradation of P_o into orthophosphate.

Effect of Drying on Metal Cation Concentrations and Relationship to P Distribution

The manures contained 7 (PM-1) and 11 (PM-2 and PM-3) g kg^{-1} of magnesium (Mg) (Table 1). Calcium (Ca) content of the manures was 153 (PM-1), 152 (PM-2), and 70 (PM-3) g kg^{-1} . These differences are probably related to across-farm differences in animal rations as well as dietary differences due to bird production status. To explore the chemistry behind drying effects on labile P change, the relationship of P to Ca and Mg content in H_2O and NaHCO_3 fractions of poultry manure were examined. Significant differences were found in the ratio of P_i to Ca in H_2O -extractable P with 22°C drying (all three manures), freeze drying (PM-1), and 65°C drying (PM-1 and PM-3) (Table 4). In addition, P_{tot} to Ca ratios in these manures roughly paralleled the results of P_i to Ca ratios, indicating that the differences were due to P_i rather than the P_o component of P_{tot} . In PM-2, the P_i and P_{tot} to Ca ratios increased with all drying methods. However, freeze and oven drying decreased the ratios of P_i and P_{tot} to Ca + Mg, thus suggesting that Mg may play a larger role in this particular manure. Similar analysis of NaHCO_3 -extractable P_i and P_{tot} revealed that P_i and P_{tot} to Ca ratios change significantly with freeze and oven drying in all three manures. Kleinman et al. (2005) found positive correlations between water-extractable P and Ca, Mg, and Fe from livestock manure. Furthermore, drying can reduce water-extractable Ca and Mg in broiler manure, with the greatest reduction seen in air- and 65°C-dried samples (Sistani et al. 2001). Phosphorus solubility in poultry manure has been proposed to be controlled by solid mineral P phases (Sistani et al.

Table 4. Molar ratio of inorganic (P_i) or total (P_{tot}) P to Ca and Mg in H_2O and $NaHCO_3$ fractions of poultry manure

Treatment	H_2O				$NaHCO_3$			
	$P_i:Ca$	$P_i:Ca + Mg$	$P_{tot}:Ca$	$P_{tot}:Ca + Mg$	$P_i:Ca$	$P_i:Ca + Mg$	$P_{tot}:Ca$	$P_{tot}:Ca + Mg$
PM-1								
Undried	0.66a ^a	0.31a	0.91a	0.42b	1.13a	0.52a	1.75a	0.81a
Freeze dry	0.39c	0.22c	0.57b	0.32c	0.62c	0.38b	0.98b	0.61b
22°C	0.59b	0.28b	0.92a	0.44a	0.90ab	0.47ab	1.29b	0.68ab
65°C	0.27d	0.17d	0.39c	0.24d	0.81bc	0.47ab	1.33b	0.77ab
PM-2								
Undried	4.72b	2.83a	5.87b	3.46a	2.64a	0.75a	3.04a	0.87a
Freeze dry	6.34b	1.80b	7.88b	2.22b	1.62c	0.60bc	1.92c	0.71bc
22°C	12.5a	2.93a	15.1a	3.46a	2.14b	0.66b	2.38b	0.73b
65°C	7.62b	1.58b	9.31b	1.94b	1.48c	0.56c	1.80c	0.67c
PM-3								
Undried	3.19b	2.02a	3.39a	2.15a	3.91a	1.04b	3.66a	0.97b
Freeze dry	2.98b	1.53b	2.98b	1.52b	3.03b	0.94b	2.99b	0.93b
22°C	3.50a	1.37c	3.09b	1.21c	2.57b	0.96b	2.41c	0.90b
65°C	2.70c	0.95d	2.47c	0.88d	2.59b	1.63a	2.32c	1.44a

^aAverages followed by the same letters within columns for each manure are not significantly different ($\alpha = 0.05$).

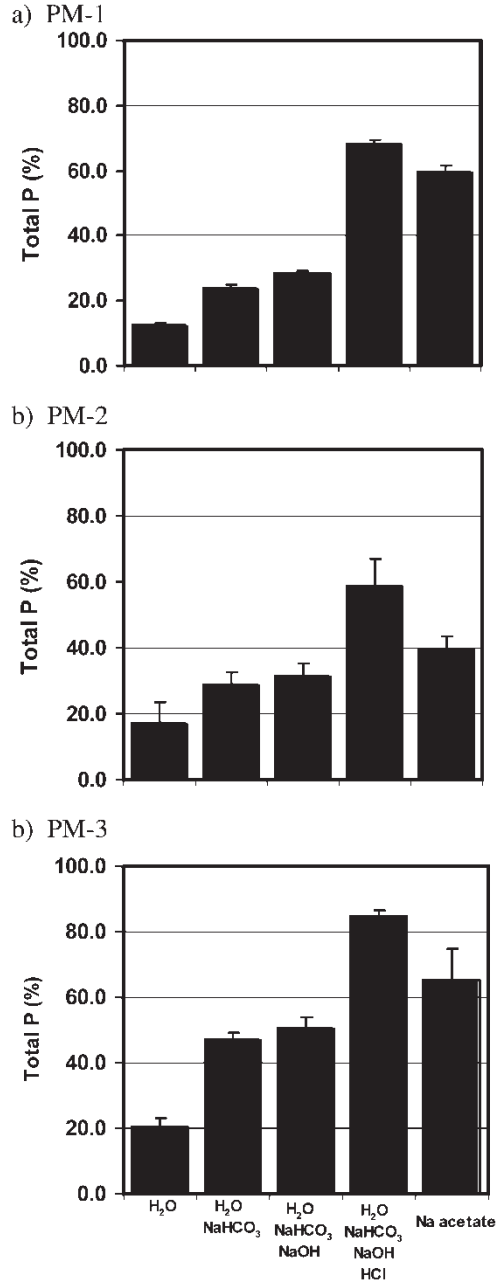


Figure 2. Comparison of P extracted by sodium acetate buffer (100 mM, pH 5.0) with that from sequential fractions of freeze-dried poultry manure. Horizontal bars represent standard deviations (n = 9).

2001). Studies by Ajiboye, Akinremi, and Racz (2004), Sharpley and Moyer (2000), and Dou et al. (2000) showed that the concentrations of Ca, Al, and Fe control P solubility in manure. Positive correlations between water-extractable Ca and P and the presence of Ca as the dominant manure cation suggest that Ca-P forms would control P solubility (Dou et al. 2000; Kleinman et al. 2005). This work supports this proposition.

Comparison of Acetate Buffer–Extracted P to That from Sequential Extractions

Sodium acetate buffer (100 mM, pH 5.0) has been tested as a single extractant to determine plant-available P in dairy manures because extraction conditions are close to those for Morgan (1.24 M sodium acetate buffer, pH 4.8) or modified Morgan P [0.62 M ammonium oxide (NH_4OH) + 1.25 M acetic acid, pH 4.8], which have been used for soil P testing (Akinremi et al. 2003; Griffin, Honeycutt, and He 2003; He, Griffin, and Honeycutt 2004b). He, Griffin, and Honeycutt (2004b) found that dairy manure P extracted by acetate buffer was equivalent to the summed amount of P removed by H_2O , NaHCO_3 , and NaOH sequential fractions. In this study, poultry manure contained small quantities of NaOH-extractable P and a relatively large amount of HCl-extractable P (Figure 1), thus providing an opportunity to verify the correlation between acetate buffer–extracted P and sequentially extracted P. In PM-1, acetate extractable P was two times higher than that extracted by H_2O , NaHCO_3 , and NaOH together but less than the total extracted by H_2O , NaHCO_3 , NaOH, and HCl (Figure 2). Similarly, acetate-extractable P was about 22% higher in PM-2 and PM-3 than H_2O , NaHCO_3 , and NaOH but less than the total extracted by H_2O , NaHCO_3 , NaOH, and HCl, suggesting that the NaOH-P fraction of animal manure is extractable by sodium acetate buffer. These observations imply that whereas sequentially extracted NaOH- and HCl-extractable P in soil are considered less plant available than H_2O and NaHCO_3 fractions (Cross and Schlesinger 1995), the counterpart P in animal manure should not be assumed to be so.

CONCLUSIONS

Using the sequential fractionation procedure, the effects of drying process on the P distribution in poultry manures was investigated. In the three manures tested, the most significant change was observed in the water-soluble P fraction. The concomitant increase in H_2O -extractable P and decrease in NaHCO_3 -extractable P with drying suggests that a transformation occurred whereby P was becoming more labile under the influence of heat associated with drying. In addition, changes were observed in NaOH-extractable P and HCl-extractable P from one manure, leading to decreases in the amount of

recalcitrant and residual P after drying. It seems that the drying process caused moderately stable P in poultry manure to either become more labile or more recalcitrant. This bilateral trend might partly explain the inconsistent observations of drying reported by other research groups. Therefore, the variations due to drying effects should be taken into consideration when evaluating manures for P availability or when comparing data in which different drying methods have been utilized.

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