Introduction

Phosphorus (P) is considered one of the most important nutrients for plant growth. It is used as a fertilizer either commercially available or in the form of manure, and also plays a key role in the growth of livestock. Animal manure of one of the sources of P. The concentration of P varies with the age of livestock, diet, and bedding type. It also depends on the livestock species. P found in animal manure exists in organic and inorganic form. In general, 45-90% of manure P is inorganic in nature and the other is supposed to be organic P [1]. Composted manure is richer in organic nutrients than fresh manure [2]. However, fresh, non-composted manure has higher nitrogen (N) content than the composted manure. Fresh manure may contain high amounts of viable weed seeds. In addition, various pathogens may be present in fresh manure, which can
cause illness unless proper precautions are taken. On the other hand, the residual effects of the manure and compost are important. Continuous use of manure or compost can lead to high levels of residual P and other nutrients, which can potentially be transported to lakes and streams as a runoff or leach to pollute the groundwater. Besides, the presence of heavy metals in composts raises serious concern about the adverse environmental impact. High and excessive accumulation of heavy metals in soil and other media contaminate both human and animal food chains [3-4]. Overuse of synthetic fertilizers has resulted in serious environmental concerns in many industrial countries, while in developing countries they are neither available nor affordable. Economic and environmental issues associated with intensive use of chemical fertilizers have thus generated an interest in alternative management systems. The application of PL waste to agricultural lands has resulted in excessive soil P in many locations. If over applied, poultry manure can be a source of P enrichment of watersheds [5]. Excess soluble P could endanger surface water quality through eutrophication [6]. To address these concerns, many countries in the world have established specific guidelines and standards for the application of composts in agricultural lands.

Because of environmental concerns, future applications of organic amendments should be based on P content. Animal manure and bedding materials contain P in organic forms. After microbial mineralization from the organic forms, the P applied to soils is subject to the same fates as inorganic fertilizer P. The application of animal wastes should be based on soil tests, nutrient content of the material, and crop needs. The phosphate in fertilizers and manure is soluble. Manure contains soluble phosphate, organic phosphate, and inorganic phosphate compounds. Agronomic practices are aimed at reducing the dependence on inorganic fertilizers for sustainable agriculture. One of the measures being adopted for relieving environmental problems arising from agricultural production is to recycle animal manure and other organic products as fertilizers [7-9]. Detailed information about the chemical composition of soil P is necessary to understand the plant nutrient and soil biogeochemical cycles as the soil P forms vary widely in bioavailability [10]. Sequential extraction procedures are used to fractionate soil inorganic and organic P according to its chemical stability [11]. Characterization of P in the animal waste may help to ascertain the behavior of P and identify sustainable field management practices to reduce possible P losses. The objectives of the present study were to compare different types of fresh and composted animal manure for the release of P fraction and to determine heavy metals (Fe, Zn, Mn, Ni, and Hg) in composted and fresh manure.

### Experimental

#### Study area

The present study was conducted in the village of Pohar in the Abbottabad District located in Hazara division of Khyber Pakhtunkhwa Province of Pakistan (Fig. 1). Total area of the district is 1,969 km² along with Abbottabad city, which is its major urban area. Approximate population of the district is 1.182 million.

#### Sampling and Analysis

Animal compost and fresh manure samples of cow, buffalo, poultry, and goat were collected, air dried, crushed, and sieved (<0.5 mm pore size). The sieved samples were digested in a mixture of duplicate acids (HNO₃ and HClO₄). Metals (Ni, Fe, Zn, and Hg) in the extract of digested manure samples, were determined by atomic absorption spectrophotometer [12]. Samples weighing 0.25 g were digested with 5 mL concentrated HClO₄ by gradual heating. After drying, 20% HNO₃ was added to the sample and it was heated again for 1 h. The solution was diluted to 50 mL with deionized water and passed through a 0.22 μm filter. The P was determined on a spectrophotometer using the phosphor molybdate blue method [13]. Absorbance was determined at a wavelength of 710 nm. The manure samples were extracted for water-soluble K, Ca, Mg, and other metals using de-ionized water. A water extract removes only the easily dissolved form of elements but very little of the adsorbed and mineral forms (K, Ca, Mg, and other metals) were determined on a polarized Zeeman atomic absorption spectrophotometer (Model Z-2300, Hitachi, Japan). Soil pH was measured in soil-water (1:5; w:v) suspensions (Table 1). Treatments were replicated thrice.

#### P Fractionation

Manure samples weighing 0.5 g were placed in a 100 mL centrifuge tube. P fractions were determined by the modified sequential extraction procedure of Hedley et al. [14], as described by Ajiboye et al. [15]. The fresh and composted manure samples weighing 0.5 g were
placed in a 50 mL centrifuge tube and were fractionated into readily plant-available P, labile inorganic P (another plant-available fraction), Al-associated inorganic P, and Ca-associated P by sequential extraction with 30 mL of de-ionized water, 0.5 M \( \text{NaHCO}_3 \) (pH 8.5), 0.1 M \( \text{NaOH} \), and 1 M \( \text{HCl} \), respectively. For each extraction, the material was shaken for 16 h at room temperature. After each end-to-end shaking the tubes were centrifuged at 10,000 rpm for 10 min., and the supernatants were filtered by a 0.2 μm filter. The shaking/filtering cycle was repeated for all the extracts. Total metals in the samples were determined with a mixture of \( \text{HClO}_4 \)-\( \text{HNO}_3 \) [12]. Samples weighing 0.25 g were digested with 5 mL concentrated \( \text{HClO}_4 \) by gradually heating it over a hot water bath for 1 h. After drying, 20% \( \text{HNO}_3 \) was added to the sample and it was heated again for 1h. The solution was diluted to 50 mL with deionized water and passed through a 0.22 μm filter. P was determined on a spectrophotometer using the phosphomolybdate blue method [13]. Absorbance was determined at a wavelength of 710 nm. Samples were replicated three times. Data were analyzed using Stat view (SAS, 1999) software [16]. Means were compared for P fractions by LSD-test at 0.05 level of significance.

### Results and Discussion

P fractionation of composted and fresh manure showed that the P forms changed significantly during fractionation. The overall sequential extraction for reagents considerably varied for P in the order \( \text{HCl} > \text{NaOH} > \text{NaHCO}_3 > \text{H}_2\text{O} \). The mean levels for the release of P obtained in water were higher in fresh and varied in the order of (PL) > (CW) > (GT) > (BF), whereas P levels in the compost were PL > CW > GT > BF. Water-soluble extracted P increased to the maximum in fresh PL (154.0 mgkg\(^{-1}\)) and the lowest was noted in BF compost (89 mgkg\(^{-1}\)) as shown in Fig. 2. Sharpley and Moyer [6] reported a reduction in water-soluble P following the composting of poultry manure, apparently due to the diluting effect of the added C sources, whereas Sikora and Enkiri [17] found no significant effect of composting on water-soluble P. The mean levels for the release of P obtained in NaHCO\(_3\) was the maximum in composted manure and varied as PL > CW > GT > BF; whereas the concentrations in fresh manure were noted as PL > CW > GT > BF (Fig. 2).

Extracted P increased to the maximum in PL compost (164.5 mgkg\(^{-1}\)) while the lowest concentration was noted in BF compost (99 mgkg\(^{-1}\)) as shown in Fig. 2. For NaOH-extracted P, nutrients were greater in composted manures than those observed in the order of PL > CW > BF > GT, whereas in fresh it was PL > CW > GT > BF, respectively, maximum concentration was observed in composted PL (99 mgkg\(^{-1}\)) and the lowest was in fresh BF (86 mgkg\(^{-1}\)). The low concentration in fresh manure could be due to the use of additives such as ferrous-ferric hydrogen sulfate and alum litter treatments. Moore et al. [18] applied alum-treated poultry litter to pastures for three years and found that soluble reactive P concentrations in runoff were 75% lower than normal litter. Concentrations of bioavailable inorganic P (NaHCO\(_3\)-IP), readily mineralizable organic P (NaHCO\(_3\)-OP), potentially bioavailable inorganic P (NaOH-IP), and potentially bioavailable organic P (NaOH-OP) were still higher in the media amended with compost because of higher total P concentration in the compost. P extracted with HCl in composted manure were PL > CW > GT > BF, while in fresh they varied in the order of PL > GT > CW > BF. A higher concentration was found in PL compost (106 mgkg\(^{-1}\)), whereas the lowest was observed in fresh BF (70 mgkg\(^{-1}\)), respectively (Fig. 2).
McDowell and Stewart studied the P concentration in fresh and dry dung of sheep, cattle, and deer. They found a higher concentration of P in sheep and reported low bioavailability and decreased P in dung with drying [19]. Williams et al. [20] found that acid P constitutes stable Ca-bound P so it would unlikely contribute to the eutrophication of water sources. However, it could be mobilized to labile fraction when the ready P is depleted. The causes of increased P availability under reduced conditions have been ascribed to the reductive dissolution of Fe oxides and the liberation of sorbed and occluded P, changes in soil pH that affect solubility of P-compounds, and the desorption of P from surfaces [21]. The relative differences observed for the NaOH-P fraction across the fertilizer treatments were essentially similar to the HCl-P form. The NaOH-P is a chemisorbed form [20]. It also contains P associated with humic compounds and P sorbed to Fe and Al oxides. The depletion of water-soluble P and NaHCO₃-P (the readily available P) suggest that these fractions of P may either be lost through leaching or runoff. The threshold agronomic P level was reported as 20 mg kg⁻¹ soil by [21]. It is also reported that aerobic-anaerobic condition changes in pH, and organic matter influence the solubility of P forms [22]. Cattle fed diets containing the highest concentration of P also produced manure with the highest concentration of total P [22], but cattle fed on high fat have no impact on manure composition [24].

Eghball and Power [25] suggested that P-based compost or manure application is most effective for corn production and increasing soil phosphorous. Takahashi [26] studies the characterization of the different forms of P from organic soil amendments by using sequential P fractionation. He reported that >60% of total P was inorganic, which in major inorganic P were NaHCO₃- and HCl-extractable P fractions in the cattle manure composts. Increasing manure concentration resulted in a linear increase in manure water-extractable P and TP. P availability from all manure types is high. The P in manure can be used similarly to P fertilizer. The elevated soil P on the count of excess manure application is an indicator of its potential transfer from the manure. All the P forms except water P were strongly related to total P in the soil. The concentration of P extracted by water significantly differed between compost and fresh poultry litter.

The measurement of water-soluble P in the manure is important because it includes available forms of P that can immediately be bioavailable and thus have the potential to contribute to environmental contamination. The environmental risk of applying organic amendments depends on their total P content as well as available P [27]. Generally, the P in the compost used in our work is 70% inorganic, which is readily available for plant absorption, microbial turnover, and soil adsorption. In the last case, when P sorption capacity reaches saturation, P can be lost by leaching or surface runoff if the amounts applied exceed crop removal. The biological portion of P cycle is controlled primarily by bacterial and fungal decomposition, in mobilization, and mineralization. It also varies with plant uptake microbial immobilization and mineralization of P. The amount of fractions was higher in both composted and fresh manure because of the fact that these wastes have been sequentially fractionated in original form and were not applied to the soil.

The P forms extracted through NaHCO₃, NaOH, and HCl were found to be higher in composted manure while H₂O-extracted P was higher in fresh manure. The environmental risks of applying organic amendments depend on their total P contents as well as forms of P [27]. Meek et al. (1979) [28] reported an increase in total and available P due to manure application. Chapuis-Lardy et al. [29] studied the effects of various treatments of cattle slurry manure on water-extractable P. They reported total readily soluble P content in slurry manures from the nine farms ranging from 1.83 to 4.06 mg/g of dry matter. A significant percentage of total P in the slurry manure arose in the water-extractable content, which was 46-71%, and the readily soluble P fractions was 24-51%. Santillan et al. [30] reported the increase of organic matter after the application of manure and high content of inorganic nitrogen and phosphorous after seven years of manure application. They suggested the monitoring of surface and well-water quality due to leaching and the prevailing application of manure as fertilizers.

Irshad et al. [31] tested the concentrations of plant nutrients like C, N, P, K, Na, and B in fresh and composted manures of buffalo, camel, cow, goat, and poultry. They reported increased extractable P and B while decreasing
concentrations of total N, C, extractable Na, and K with composting. Similarly, Li et al. [32] conducted a comparative study of characterizing P in dairy, animal manure, swine, and broilers. They reported that availability, quantity, and P form varied between different animal manures. Gagnon et al. [33] studies the different forms of P both in compost and compost-amended soils. They reported that inorganic P was the most significant form of P both in the compost and compost-amended soils. Another study reported that P forms from poultry, cattle, and swine dry manures are bioavailable and in organic in nature and dependent on Ca:P ratio in manures [34]. Ajiboye et al. [35] reported 30 to 40% of total hog and cattle manure P and 10% of total biosolids P was extracted with water, and 21 to 32% of total P in all organic amendments was extracted with NaHCO₃ except in dairy cattle manure (45% of total P).

The manure practices appear to control the chemical forms of heavy metals. The metal contents varied in the order Fe > Mn > Hg > Ni > Zn (Fig. 3). The metals in composted manure were observed as Fe > Mn > Hg > Ni > Zn and in fresh manure appeared in the order Fe > Mn > Ni > Hg > Zn. A higher concentration of Fe was found in BF compost with (338 mg kg⁻¹) and the lowest in fresh PL (111 mg kg⁻¹) as shown Fig. 3. The maximum concentration of Zn was found in BF compost (10.35 mg kg⁻¹) and the minimum were in fresh PL (2.3 mg kg⁻¹). A high concentration of Mn was found in BF compost (71.3 mg kg⁻¹) and low were observed in fresh PL (38 mg kg⁻¹) as shown in Fig. 3. The higher concentration of Ni was observed in BF compost (20.8 mg kg⁻¹) and the lowest in fresh PL (14 mg kg⁻¹) as shown in Fig. 3. The maximum concentration of Hg was noted in PL compost (37.5 mg kg⁻¹) and minimum in CW fresh (6.2 mg kg⁻¹) as shown in Fig. 3. Faridullah et al. [36] compared the various trace elements in both fresh and composted animal manures by adopting sequential fractionation procedure. They used both fresh and composted manures from four different species of animals like buffalo, cow, goat, and poultry to find fractions of iron, nickel, zinc, mercury, and manganese. They found that Fe, Mn, and Hg were in higher concentrations in composted samples, while Zn and Ni were mainly higher in fresh manures. Concentrations of heavy metals in plants are affected by sheep manure and inorganic fertilizer, and especially Pb, Cd, and Zn concentrations are decreased by the applications of sheep manure in plant tissues [37].

Bolan et al. [9] reported that the concentration of metals in manure can vary considerably depending on the number of factors, including type of livestock (i.e., poultry, swine, or cattle). Within a type, additional variation is associated with the age of the animal, type of ration, housing type, and waste management practice. The metal contents of manure by-product depend primarily on the amount used in the feed and health remedies. Increases in metal concentration in animal feed have often resulted in corresponding increases in their concentrations in the manure by-products [38]. Heavy metals were investigated in 109 samples in Korea and reported that Zn and Cu were found at their highest concentrations in compost other than the tested metals [39].

DeLaune and Moore [40] studied the Cu and Zn concentrations, which were measured in runoff water collected from plots fertilized with composted and fresh poultry litter and in which were found elevated concentrations of Cu and Zn in compost as compared to poultry litter. Another study reported a high concentration of heavy metals in solid municipal waste composting due to uncontrolled leaching, along with runoff and compost, which subsequently degraded water quality and elevated environmental risk [41]. Eneji et al. [6] reported that the major concern of soil pollution with available heavy metals by the use of manures can be minimized through composting under aerobic conditions. The bacterial community plays an important role, especially phosphate-solubilizing bacteria. Wei et al. [42] suggested the combined use of bacteria and biochar as an effective strategy for increasing available P fraction as both these have important roles in transformation. Vazquez et al. [43] studied the physicochemical quality of compost and reported that compost quality is not affected by moisture (80%). Pare et al. [44] reported a decrease in the extractability of some heavy metals during composting of biosolids and municipal solid wastes, indicating a reduced risk of their entering the food chain through crops and water. According to Pare et al. [44], proportionally greater increases in humic compounds during composting will cause more heavy metals to interact with sites of greater complexing strengths, where they become less accessible and available to plants and extracting agents.
According to scientific risk assessment, the presence of cadmium in P-containing fertilizer does not pose any risk to human health [45]. Tian et al. [46] suggested the Caragana microphylla-straw compost as an alternate for excessively fertilized soils, and for reducing environmental risk because excessively fertilized soil releases more nutrients. Phosphorus loss risk coefficient was studied by Robert and reported that soil type, manure type, and their interaction has a significant effect on P loss risk coefficient, which is important for assessing phosphorus and nutrient management in the environment [47]. Some microbes play an important role in increasing soil fertility along with plant growth, but some work in depleting P and compost clearing [48-49].

Phosphorus absorption by soil is highly important for management practices, depending on different transport factors like soil hydrological conductivity, precipitation, slope, and distance, along with integration, which plays an important role in P absorption [50]. Leaching of P and other elements from manure to water sources may affect surface water quality [51-52]. Vermicomposting can play an important role in stabilizing heavy metals in composting animal manures [53]. This can help in achieving finer homogeneity than for a conventional product [54]. Animal manure compost as fertilizer not only is important for crops and plants, but also for cultivating microalgae for the production of different bio-products and waste management, as it is a rich source of nutrients [55]. It is suggested in some studies that metals may be cautiously inspected in both fresh and composted manure before applying it to agriculture [56-57], while in others composting is one of the treatment technologies for solid waste management [58-60].

Conclusions

The overall sequential extraction for reagents considerably varied for P in the order of HCL > NaOH > NaHCO₃ > H₂O. The P and other nutrients availability increased progressively during composting with the simultaneously decrease in fresh manures. The metals were varied in the order Fe > Hg > Mn > Zn > Ni because the type of bedding material in manures may influence its chemical properties. Our results demonstrate that agronomically and environmentally important attributes associated with composted and non-composted manure application can be predicted based on P fractions and an optimization procedure involving regression analysis. Composting animal manures before land application can affect the forms and concentrations of P. The use of composted animal manures as soil amendment could be economically productive and would potentially ameliorate the major concerns associated with animal waste and acid soil management.

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