Alternative means of recycling pineapple leaf residues

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An alternative means of recycling pineapple leaf residues.

Abstract — Introduction. One of the challenges of the Malaysian pineapple industry is to develop new techniques for managing pineapple residues. A study was carried out to investigate whether K-humate can be produced from these residues. Materials and methods. Pineapple leaves were air-dried, shredded and chipped. Some of the shredded leaves were incinerated at 500 °C. Potassium hydroxide (KOH) was extracted by dissolving the ash obtained in distilled water for 24 h at a ratio of 1:7 (ash / water) after which the sample was filtered. After this extraction of KOH from the ashes of pineapple leaf residues, the remaining ash residues were further analyzed to assess the total and soluble K contents. Moreover, a given quantity of shredded leaf residues was mixed with chicken dung (as a source of microorganisms and nitrogen) and chicken feed. The mixture was composted with standard procedures. Humic acids were reconstituted using KOH from pineapple leaves with K from ash residues and K from composted pineapple leaves. Results and discussion. By reconstituting humic acids with KOH, a K-humate was produced with approximately 34.5% of its K readily soluble in water. A reconstitution of humic acids with K from ash residues produced a K-humate with approximately 3.34% of its K readily soluble in water. Conclusion. The K-humate produced from the reconstitution of humic acids with K from KOH can be used in fertigation programmes as a source of K while the K-humate produced through the reconstitution of humic acids with K from ash residues can be used as a source of K for fresh water fishes.

Malaysia / Ananas comosus / crop residues / leaves / composts / potassium hydroxide / potassium / humic acids

Réutilisation des résidus de feuille d’ananas.


Malaisie / Ananas comosus / résidu de récolte / feuille / compost / hydroxyde de potassium / potassium / acide humique
1. Introduction

In Malaysia, about 13 t·ha⁻¹ of pineapple residues are produced on peat each growing season; without an effective and efficient way of handling them, these residues are simply burnt before planting the next crop. Open burning not only reduces the quantity of the valuable organic matter added to the soil but also produces a large quantity of gases such as carbon dioxide which are implicated in ozone depletion. In 1997, fires and haze outbreak across Southeast Asia cost Malaysia about 321 million US$ [1]. Furthermore, even though as much as 1.31 t·ha⁻¹ ash with a potassium concentration of 17% is produced from burning pineapple residues, this practice increases neither the uptake of K significantly nor the crop yield [2]. This has been attributed to a low recovery of K since, besides early application before planting through burning as compared with fertilizer application 65 d after planting, the K uptake in pineapple is generally slow during the early growth period [3]. Thus, due to high rainfall, K is lost through leaching [2].

One of the challenges of the Malaysian pineapple industry is to develop new techniques for managing pineapple residues. Therefore, the ever-increasing awareness about environmental pollution, the need for achieving sustainable agricultural development and the continuous increase in fertilizer price call for alternative techniques for managing this valuable resource. One of the possible ways of achieving this objective is to develop value-added product(s) from pineapple residues. This can be achieved by producing potassium hydroxide (KOH) from pineapple leaf residues, which, in turn, can be used for extracting humic acid from composted pineapple leaf residues for the production of a potassium-rich humic acid called K-humate.

Our study was carried out to investigate whether K-humate can be produced from pineapple leaf residues by reconstituting humic acid extracted from composted pineapple leaf residues using KOH (from pineapple leaves) with K from ash residues (after extraction of KOH from leaves) and K from KOH from pineapple leaf residues, respectively.

2. Materials and methods

Pineapple leaf residues were collected from the Simpang Rengam Pineapple Estate, Simpang Rengam, Johor. The leaves were air-dried, shredded and chipped using the Briggs and Stratton shredder-chipper (Model 135212). Some of the shredded leaves were incinerated at (300, 350, 400, 450 and 500) °C for 4 h in a muffle furnace but, because 500 °C produced the best ash from the shredded leaves (almost white), this ash at 500 °C was used throughout this study. The ash was digested using the dry ashing (single dry ashing) method. The K concentration in solution was determined using an atomic absorption spectrophotometer (AAS).

The KOH was extracted by dissolving ash in distilled water for 24 h at a ratio of 1:7 (ash / water, weight / volume basis), after which the sample was filtered through a Whatman filter paper number 2. This ratio was arrived at by analyzing the molarities of a series of ash (shredded leaf residue incinerated at 500 °C) to distilled water ratios of 1:5, 1:6, 1:7, 1:8, 1:9, and 1:10. The ash to distilled water ratio of 1:7 was chosen because the ratio produced approximately a 0.1 M KOH solution for the extraction of humic substances [4]. The method of acid-base titration was used to confirm this concentration. The 0.1 M KOH solution obtained (not the acidified one) was analyzed for K, Ca, Mg, Na, Cu, Zn, Al, Fe and Mn using an AAS, and pH assessed with a pH meter.

After the extraction of KOH from the ashes of pineapple leaf residues, the remaining ash residues were further analyzed to assess the total and soluble K contents. Ash residues were oven-dried at 60 °C until a constant weight was attained. Some of the ash residues were digested for the total K using the single dry ashing method. For soluble K, we used the method described by Bailey et al. [5].
A quantity of 25 kg of shredded leaf residues was mixed with 1.25 kg of chicken dung in liquid form (as a source of microorganisms and nitrogen) and 1.25 kg of chicken feed. The mixture was composted in a composting drum for 28 d. The moisture content of the compost was maintained at (55 to 60)%. The ambient temperature and the compost temperature were measured on a daily basis using a thermometer. The total organic carbon (TOC) of composted and uncomposted pineapple leaf residues was measured by loss of weight on ignition at 550 °C for 8 h (organic C calculated as 58% of the organic matter) [6]. The micro Kjedahl method was used for the determination of the total nitrogen [7] and the leaching method for the determination of the cation exchange capacity (CEC) [8] of the composted and uncomposted leaves. For the extraction and purification of humic acid, the method described by Stevenson [9] was followed.

### Table I.
Elemental composition of a 0.1 M KOH solution (50% K) obtained from pineapple leaf residues.

<table>
<thead>
<tr>
<th>Element</th>
<th>Content in the KOH solution (mg kg⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Al</td>
<td>2</td>
</tr>
<tr>
<td>Cu</td>
<td>17</td>
</tr>
<tr>
<td>Mn</td>
<td>30</td>
</tr>
<tr>
<td>Zn</td>
<td>40</td>
</tr>
<tr>
<td>Fe</td>
<td>80</td>
</tr>
<tr>
<td>Mg</td>
<td>120</td>
</tr>
<tr>
<td>Ca</td>
<td>300</td>
</tr>
<tr>
<td>Na</td>
<td>1300</td>
</tr>
</tbody>
</table>

3. Results and discussion

The total K concentration in the pineapple leaf residues was high (35%) and was sufficient to produce a 0.1 M KOH solution. This high K concentration may be due to the naturally high K content in pineapple leaves [2, 3]. By incinerating the pineapple leaf residues, a unit weight of pineapple leaf residue ash will naturally be high in K.

The pH of the 0.1 M KOH solution was 11.10. The 0.1 M KOH solution contained 50% K with Al, Cu, Mn, Zn, Fe, Mg, Ca and Na content generally low (table I). From a practical point of view, further purification of the KOH was not deemed necessary as this would entail some cost which would in turn make the production of this KOH more expensive. This could possibly be one of the reasons why an analytical grade of NaOH is preferred to that of an analytical grade of KOH in humic and fulvic acid extraction. Aside from this, the elements in the KOH solution obtained from the pineapple leaf residues could be recycled in the form of fulvate, particularly in K-fulvate, after the extraction of humic acid or K-humate (salt of humic acid).

During the composting process, three typical phases of composting were observed by temperature measurements (figure 1). There was a mesophilic phase during the first 2 d in which the temperature rose to 46 °C. At this mesophilic stage, the predominant microbes are the mesophilic bacteria [10]. At this time, the abundance of readily available and easily digestible substrate (e.g., sugars, starches, simple protein compounds, etc.) might have ensured that the microorganisms were very active, leading to the generation of large quantities of metabolic heat energy, which caused the temperature of the compost to increase.

### Figure 1.
Ambient and compost temperature measured at different times during the composting of pineapple leaf residues.
The second stage was a thermophilic phase lasting for 15 d in which the temperature increased to a maximum of 67.3 °C. As the temperature rose past 46 °C, conditions were less favorable for the mesophilic bacteria and instead began to favor the thermophilic bacteria, mostly *Bacillus* species [11] that play a major role in the decomposition of proteins and other carbohydrate compounds [12]. The resulting increased microbial activity of the thermophiles caused the temperature in the compost to rise to 67.3 °C. Further temperature increases were avoided by turning and, mixing the compost once every 2 d, and after 28 d of composting, the compost temperature equaled that of the ambient temperature.

Eventually, with the depletion of the food sources, overall microbial activity decreased and the temperature fell, resulting in a second mesophilic phase during the cooling stage. As the readily available microbial food supply was consumed, the temperature fell to ambient and the material entered a maturation stage and, during this stage, the microbial activity was low [10]. The phases described were similar to the results previously reported by several researchers during the composting process [6, 13–16].

The change in CEC [17], ash content [6] and the C/N ratio [6, 14, 18] reflects organic matter decomposition and stabilization of a compost. The ash content increased from the initial value of 7% for the uncomposted leaves to 23% for the composted leaves. The CEC of the compost, a measure of the capacity of the compost to hold exchangeable cations such as K, Ca, Mg and Na to negatively charged surfaces of the compost (e.g., OH, COOH, when dissociated) also increased from an initial value of (32 to 68.32) cmol·kg⁻¹. The high CEC of the compost indicates that the organic material of the compost has become more humified [19]. The carbon to nitrogen (C/N, weight/weight basis) ratio decreased from an initial value of 38.30 to a final value of 11.58. The [C/N] ratio of 11.58 found for the composted pineapple leaf residues in this study compared well with the ratio of 12 reported by Chefetz et al. [6]. The ratio also fell within the [C/N] ratio of 10 to 12 usually considered to be an indicator of stable and decomposed organic matter [14]. The humic acid increased from 2.77% in the uncomposted leaf residues to 20% in the composted leaf residue.

The total and soluble K of the ash residue were (18 and 8.34)%, respectively. There was a negative significant relationship between the amount of soluble K released from the ash residue (after extraction of KOH) with equilibration time (figure 2). The highest release occurred a day after equilibration followed by a successive decrease with time, indicating that the highest amount of soluble K that can be released from the ash residue after extraction of 0.1 M KOH occurs within a day and does not exceed approximately 8.34% K even if allowed to stand for one week.

In order to make use of this ash residue, humic acid was reconstituted with K from the ash residue using a humic acid to ash residue ratio of 10:1 (weight basis), and equilibrated in 40 mL distilled water. There was a significant negative relationship between the amount of K adsorbed at the exchange complex (perhaps COOH, phenolic OH, etc.) of the humic acid with time (figure 3). A day after equilibration in distilled water, there was a general decrease in the adsorption of K, indicating that the maximum adsorption does not go beyond 1 d. It seems that the general decline in the soluble K of the ash residue observed in figure 2 partly explains this observation, as a decrease in soluble K led to a decrease in the amount of K adsorption, and this association was found to be significant (figure 4). It is interesting to note that
Recycling pineapple leaf residues

approximately 3.34% K was in the humic acid, and this amount is sufficient to support the K requirement of fresh water fishes.

Fresh water contains less than 10 mg K-L⁻¹ [20]. According to the studies of Shearer [21], inadequate levels of dietary K supply for the young king salmon (a fresh water fish) resulted in poor feed intake and feed utilization as well as growth retardation when the diet contained less than 1 mg K-L⁻¹, and increased mortality also occurred. A maximum growth was obtained at a dietary level of 8 g K-kg⁻¹ dry diet, while whole-body concentrations were normal at 6–12 g K-kg⁻¹ diet [21]. Besides, the backbone of K-humate, which is humic acid, can stimulate the growth of organisms such as phytoplanktons and algae [22], diatoms and dinoflagellates [23]. These organisms play an important role in the food chain of fishes. It is believed that humic acids stimulate the growth of these organisms by chelating and solubilizing micronutrient metal ions that otherwise would be present in an insoluble or colloidal form. Furthermore, N and P directly from humic acids enhance growth of these organisms [23]. Excessive levels of Cu are toxic to fishes [24] and a study conducted by Florence et al. [25] has indicated that humic substances may be effective in ameliorating the toxicity of hydrophobic Cu complexes in fresh water.

The relatively low amount of K displaced (3.34%) may be partly attributed to the low acid strength (pKa = 15.7) of water which in this case served as the conjugate acid of the humic acid salt (K-humate) [26]. Almost all the K in the salt seemed to have been displaced from the exchange sites (carboxylic and phenol) [26] when the salt was acidified with dilute HCl before purification to obtain humic acid (figure 5) for the onward reconstitution with K.

One of the variables that is used to assess the quality of fresh waters for fresh water fishes is pH, and the available literature has shown that waters with pH near neutral (7 ± 1) [27] or a pH range of 6.5 to 9 are likely to be suited to good fish production [28]. The pH of the K-humate was 6.7, which is consistent with the pH of fresh waters that support fresh water fish production.

In terms of a K source for agricultural use, the amount of K in the reconstituted humic acid, as noted in the preceding discussion, is relatively low. As a means of increasing the K content in humic acids, purified humic acids were reconstituted with K using humic acid to 0.1 M KOH (derived from an incinerated pineapple leaf residue) ratios (weight to volume basis) of 1:5, 1:10, 1:15, 1:20, 1:25 and 1:30 (table II). Ratios below 1:5 were excluded because of the lack of complete dissolution of humic acid in the KOH. The adsorption of K at the

![Figure 3](image3.png)

**Figure 3.** Absorption of soluble K (SK) release by humic acids, from ash residue remaining after extraction of KOH from pineapple leaf residues.

![Figure 4](image4.png)

**Figure 4.** Relationship between soluble K released from ash residue remaining after extraction of KOH from pineapple leaf residues and absorption by humic acid.

![Figure 5](image5.png)

**Figure 5.** Effect of washing on K removal from humic acid extracted from composted pineapple leaf residues.
exchange sites of the humic acids generally increased with the increasing ratios but it leveled off after 1:20, indicating that the succeeding ratios contained a sufficient amount of K to saturate the exchange sites of the humic acids with. A similar trend was observed for pH except that the leveling off started at 1:20 (Table II), probably due to the presence of the inherent cations such as Na, Ca and Mg in the KOH. Considering the amount of K in the K-humate and the pH, coupled with the fact that the K-humate readily dissolves in water, the possibility of using liquid K-humate as a source of K for fertigation or hydroponics is worth detailed exploration.

A sub-irrigation study was conducted on a poor quality soil formed on a lithological substrate of loam; irrigation water treated with humic acid extracted from sewerage sludge compost revealed that, besides increasing plant yield, the addition of humic substances stimulated the absorption of nitrogen and potassium by lucerne plants [29]. This means that, in a fertigation program, apart from serving as potassium carrier, the backbone of K-humate, which in this case is humic acid, could stimulate or facilitate the absorption of K.

4. Conclusion

K-humates can be produced from pineapple leaf residue for agricultural use.

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References


Table II.
Potassium content and pH of humic acid (K-humate) reconstituted with a 0.1 M KOH solution produced from pineapple leaf residues using different humic acid to 0.1 M KOH solution ratios.

<table>
<thead>
<tr>
<th>K-humate</th>
<th>K (%)</th>
<th>pH</th>
</tr>
</thead>
<tbody>
<tr>
<td>Humic acid only</td>
<td>Trace</td>
<td>2.20</td>
</tr>
<tr>
<td>1:5</td>
<td>25</td>
<td>9.80</td>
</tr>
<tr>
<td>1:10</td>
<td>32.15</td>
<td>10.33</td>
</tr>
<tr>
<td>1:15</td>
<td>32.95</td>
<td>10.53</td>
</tr>
<tr>
<td>1:20</td>
<td>34.54</td>
<td>10.80</td>
</tr>
<tr>
<td>1:25</td>
<td>38.10</td>
<td>10.80</td>
</tr>
<tr>
<td>1:30</td>
<td>38.12</td>
<td>10.82</td>
</tr>
</tbody>
</table>


Reutilización de los residuos de hoja de piña.

Resumen — Introducción. En Malasia, uno de los retos de la industria de la piña es desarrollar nuevas técnicas para el manejo de los residuos de este cultivo. Se efectuó un estudio para estudiar si se podía producir, a partir de estos residuos, ácido húmico rico en potasio (K-humato o humato de potasio). Material y métodos. Se secaron algunas hojas de piña al aire, se despedazaron y, luego, desmenuzaron. Algunas de estas hojas despedazadas fueron incineradas a 500 °C. Se extrajo hidróxido de potasio (KOH) disolviendo la ceniza obtenida en agua destilada (relación de 1:7, ceniza : agua) durante 24 h y, posteriormente, se filtró la muestra. Tras esta extracción de KOH a partir de residuos incinerados de hoja de piña, los residuos de ceniza restantes se analizaron para evaluar el contenido en potasio total y soluble. Por otra parte, se mezcló una cantidad dada de residuos de hojas despedazadas con estiércol de gallina (como fuente de microorganismos y nitrógeno) y con comida para aves. La mezcla se compostó utilizando procedimientos estándar. Se reconstituyeron ácidos húmicos a partir del KOH procedente de las hojas de piña utilizando el K procedente de los residuos de ceniza y el K procedente de las hojas de piña compostadas. Resultados y discusión. Al reconstituir ácidos húmicos a partir del KOH, se obtuvo ácido húmico rico en potasio con aproximadamente un 34,5% de su K fácilmente soluble en el agua. Una reconstitución de ácidos húmicos con K procedente de los residuos de ceniza produjo un “K-humato” con aproximadamente el 3,34% de su K fácilmente soluble en agua. Conclusión. El “K-humato” producido a partir de la reconstitución de ácidos húmicos con K procedente de KOH puede emplearse en programas en los que el potasio deba aportarse en fertigación, mientras que el “K-humato” producido por reconstitución de ácidos húmicos con K procedente de ceniza de residuos puede emplearse como fuente de K para pescados de agua dulce.

Malasia / Ananas comosus / residuos de cosechas / hojas / compost / hidróxido de potasio / potasio / ácido húmico