A New Method of Compost Preparation from Vegetable Waste and Dried Leaf Litters

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Abstract

Compost is decayed organic material which can be used as a fertilizer for growing plants. There are several conventional methods of compost preparation are being practiced across the country. NIPHM used the barrel composting and deep bed composting methods. For barrel composting the vegetable waste were collected from NIPHM hostel and residential staff quarters regularly. In deep bed composting method, the litters collected at quarters at regular intervals were used. To accelerate the process of decomposition bioinoculum developed by NIPHM was used. Bioinoculum and compost prepared were subjected to physicochemical and nutritional analysis. On physicochemical analysis the bioinoculum were identified as Pseudomonas sp and Bacillus sp. In NIPHM bio-inoculum method Pseudomonas sp were used as decomposer for composting called NIPHM bioinoculum method. Compost prepared by new methods undergone for physicochemical analysis i.e. moisture content, pH, electrical conductivity, and organic carbon was found as good as in control (A regular method country wide). In nutritional analysis Potassium, Phosphorous other micronutrients were found to be more than required in both the composts prepared by NIPHM and also in control. The development of composting took place in 40-50 days for barrel composting, whereas 60-70 days for conventional method. In deep bed composting, the entire process of natural composting took 90 to 120 days, whereas NIPHM modified technique took 90 days. Total 595 kg vegetable wastes were collected and 160 kg compost was harvested from all the three methods. The harvested compost was used in plot in which Spinach (Spinacia oleracea) was grown. The spinach growth in vegetable compost was compared with FYM, Earthworm compost and control (without compost). The growth of the spinach was superior in producing more root length and shoot length in vegetable compost.

Keywords
Vegetable waste, Dried leaf litters, Compost, Bioinoculum

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Introduction

In India it is estimated that nearly 70 million ton organic waste is generated annually which is either burned or land filled (Bhiday, 1994). Use of microbial inoculum to convert vegetable waste into compost is a feasible and potential technology. It is a simple biotechnological process of composting, in which certain species of bacteria are used to
enhance the process of waste conversion and produce a better end product. Many fruits and vegetables present nearly ideal conditions for the survival and growth of many types of microorganisms. In the present study, a simple microbiological process, this could provide a solution to the problem of vegetable kitchen waste disposal for recycling of solid waste into useful compost by the action of decomposing bacteria was carried out.

**Material and Methods**

**Quantity of waste generation and collection at NIPHM:**

About 30 to 40 kg of wastes are regularly generated in every month at NIPHM canteen, and NIPHM residential premises. Every house was provided with two dust bins for segregation of dry and wet wastes. The wastes are collected and cut into small pieces and transferred to plastic barrel for decomposition. Total vegetable waste collected were dried uniformly, For the collection of other wastes viz., plastic, paper, hazardous, dry, NIPHM have arranged the different bins for segregation of different wastes like plastic, paper, hazardous, dry etc. which are collected by GHMC every week. Following two treatment methods of waste management practices was developed at NIPHM.

**Development of bio inoculum at NIPHM**

To collect bioagents from the waste vegetables soil samples along with vegetable waste were collected from area where the market vegetables dumped. The soils and vegetable waste mixed samples were collected in sterile polythene zip lock covers and stored in the refrigerator without losing moisture content. Then the soil samples along with waste were rinsed thoroughly with distilled water and serially diluted up to $10^{-7}$. The highest dilutions were taken for analyzing the total microbial count by using Nutrient agar medium at 33-35°C for 24 hours.

**Barrel composting by NIPHM method**

Daily wet/vegetable waste from the NIPHM residents and NIPHM canteen were collected and then cut into small pieces and transferred to plastic barrel (100 lit capacities) on regular basis. Before pouring in to drums the vegetable pieces were mixed with coco peat, sanitizer and bio inoculum added to accelerate the composting process. Then regular mixing carried out periodically to accelerate the decomposition of wastes. The flow chart for preparation is provided in Fig.1.

**Barrel composting by Regular method**

The experiments were conducted in plastic drums of 100 lit capacities as described above. Routine method of vegetable composting was used as control treatment as check. The flow chart for preparation is provided in Fig.2. Daily 5 kilogram of the vegetable wastes were transferred into plastic barrel. NIPHM bio-inoculum 10 ml of Pseudomonas sp (pure culture) added into the wastes. In regular method the bio-inoculum was not added. Both experimental setups were periodically mixed well. Excess water will get rid of through the holes provided at the bottom of barrel. After 60-70 days compost was harvested. The composts prepared by NIPHM and Regular method were subjected to microbial and physio chemical analysis.

**Dry leaf litters wastes composting by using NIPHM bio-inoculum**

Dry leaf litters at NIPHM quarters was collected every week. This litter was converted into the compost by using the NIPHM bioinoculum. Pit method was used
for the composting and flow chart of compost preparation is provided below. The collected composts were subjected to microbial and physiochemical analysis.

**Physiochemical analysis**

Moisture Content, pH and organic carbon were determined at 0, 10, 20, 30, 40, 50, 60 and 70 days during preparation of composting. pH was determined by method described by ISI Bulletin (1982). The organic carbon was determined by the empirical method followed by Walkely and Black (1934). Moisture % was calculated (Thiruppathiet.al; 2005) for each of the compost by

a) Weighing a small container  
b) Weighing 10 g of the material into the container  
c) Drying the sample for 24 hours in a 105-110 degree C oven  
d) Re-weight the sample, subtract the weight of the container, and determine the moisture content using the following equation:

\[ M_n = \frac{(W_w - W_d)}{W_w} \times 100 \]

where:  
- \( M_n \) = moisture content (%) of material n  
- \( W_w \) = wet weight of the sample, and  
- \( W_d \) = weight of the sample after drying.

The final compost products were again tested for the pH, electrical conductivity organic carbon, Calcium, Potassium, Phosphorus and Micronutrients with a help of Soil testing laboratory, Rajendranagar, Government of Telangana.

**Results and Discussion**

**Development of Bioinoculum**

According to Bergey’s Manual of Determinative Bacteriology, the microorganisms were isolated by using King A and Kings B medium. Morphological and culture characteristics such as abundance of growth, pigmentation, optical characteristics, form, size, margin and elevation of the microbes were studied on Nutrient agar plates. The highest dilutions were taken for analyzing the total microbial count by using Nutrient agar medium at 33-35°C for 24 hours. Standard Plate Count (SPC) was carried out by spread plate Technique. Fig 5.Identification of Bacteria Gram’s staining technique was carried out to identify gram positive and gram negative bacteria. Depending upon the morphological and biochemical characters isolates were identified as *Bacillus species* and *Pseudomonas species* (Table 1).

**Physical and chemical analysis of Bioinoculum**

Twenty four hr old culture was used for the physicochemical analysis. In physical analysis it was noticed that both culture bacterial cells are rod shaped. Under the biochemical analysis 12 tests were conducted. Based on the biochemical analysis, the isolates were identified as *Bacillus* sp. and *Pseudomonas* sp. Details of Physicochemical analysis are shown in Table.2.

**Physiochemical Analysis of the compost**

The pH of the compost was lower in all the treatments than their initial values (Table 2). The decrease in pH value at the final stage of compost formation may be due to the production of CO₂ and organic acids by microbial metabolism during decomposition of different substrates in the vegetable waste (Albanell et al., 1998). Decrease in pH may be an important factor in Nitrogen retention as this element is lost as volatile ammonia at highest pH (Gautham et al., 2010). According to Viel et al., (1987) loss in organic carbon might be responsible for nitrogen
enhancement. *Pseudomonas* bacteria also have great impact on nitrogen transformation in manure, by enhancing nitrogen mineralization, so that mineral nitrogen may be retained in the nitrate form (Atiyeh et al., 2000b). Nitrogen was found high in vegetable waste compost and leaf litter compost compared to control. All other micro and micro nutrients are medium to high except Fe and Mn. In the present study, the vegetable and leaf litter wastes were effectively decomposed by the microbes. It is due to the increased microbial activity in the compost. Daywise details of the moisture content, pH and Organic carbon content day wise is provided in Table 2. Biochemical analysis was carried out for the final product also and details are provided in Table 3.

**Flow chart of Compost procedure from the leaf litters**

1. Make a pit of 2’deep X 3’width X 8’ length
2. Spread dry grass at bottom
3. Spread litters in layers on it
4. After every two layers add 1-2 kg neem leaves (dry/fresh)
5. Spray NIPHM Decomposer 10-15 ml by mixing in 1000 ml water in each layer and wet the leaves
6. Continue the process till pit is full
7. Collect the litters in polythene cover and closed it.
8. After 12 days remove the polythene cover and do mixing and again cover with polythene (Check for moisture at the time of every mixing)
9. Continue process every month till 60 days
10. Observed for decomposition state if required continue for another one month
Table.1 Morphological & biochemical profile of the bacterial isolates

| S.No | Characteristic/ Test | Isolate 1 (Bacillus Species) | Isolate 2 (Pseudomonas Species) |
|------|----------------------|-----------------------------|-------------------------------|
| 1.   | Shape                | Rod                         | Rod                           |
| 2.   | Gram’s staining      | +                           | -                             |
| 3.   | Motility             | +                           | -                             |
| 4.   | Endospore            | +                           | -                             |
| 5.   | Indole test          | -                           | -                             |
| 6.   | Methyl red test      | -                           | -                             |
| 8.   | Gelatin hydrolysis   | +                           | -                             |
| 9.   | Carbohydrate utilization | +                       | +                             |
| 10.  | Catalase test        | +                           | +                             |
| 11.  | Oxidase test         | +                           | +                             |
| 12.  | Nitrate reduction test | +                        | +                             |
| 13.  | Starch hydrolysis test | +                       | -                             |

+ Means test is positive test, - Means test is negative

Table.2 Physiochemical analysis of the compost prepared

| Days | Moisture content (%) | pH values | Organic carbon content (%) |
|------|----------------------|-----------|---------------------------|
|      | Regular Method (Control) | NIPHM Vegetable waste | NIPHM leaf litter waste | NIPHM Vegetable waste | NIPHM leaf litter waste | NIPHM Vegetable waste | NIPHM leaf litter waste |
| 0    | 58                   | 58        | 58                        | 6.5                     | 6.5                     | 6.5                     | 20                      | 22.00                  | 20.00                  |
| 10   | 56                   | 59        | 57                        | 6.6                     | 6.6                     | 6.7                     | 19                      | 22.00                  | 20.00                  |
| 20   | 50                   | 50        | 45                        | 6.8                     | 6.7                     | 6.8                     | 18.5                    | 20.00                  | 19.50                  |
| 30   | 46                   | 47        | 39                        | 6.7                     | 6.8                     | 7.0                     | 17.32                   | 19.50                  | 19.00                  |
| 40   | 41                   | 45        | 34                        | 6.9                     | 7.4                     | 7.8                     | 18.00                   | 19.00                  | 18.00                  |
| 50   | 39                   | 38        | 31                        | 7.2                     | 7.6                     | 8.0                     | 17.00                   | 18.00                  | 17.50                  |
| 60   | 38                   | 38        | 31                        | 8.0                     | 7.6                     | 8.2                     | 16.00                   | 17.5                  | 16.00                  |
| 70   | 36                   | 37        | 30                        | 8.0                     | 7.8                     | 8.5                     | 16.50                   | 15.5                 | 14.5                   |
Table 3 Chemical properties of final compost product

| Chemical Properties | NIPHM leaf litter waste compost | NIPHM Vegetable waste compost | Regular (Control) | Method |
|---------------------|---------------------------------|-------------------------------|-------------------|--------|
| Electrical Conductivity | 0.11 (Normal)                  | 0.12(Normal)                  | 0.19(Normal)      |        |
| pH                  | 8.26(Medium alkaline)          | 8.36(Medium alkaline)         | 8.2(Medium alkaline) |        |
| Organic Carbon      | Low                             | Low                           | Low               |        |
| Available Phosphorous | 14 (Medium)                    | 11 (Medium)                   | 16 (Medium)       |        |
| Available Potassium | 148 (High)                     | 107 (Medium)                  | 184 (High)        |        |
| Zn                  | 1.139 (High)                   | 0.530 (Low)                   | 0.830 (High)      |        |
| Mn                  | 0.322 (Low)                    | 0.268 (Low)                   | 0.540 (Low)       |        |
| Fe                  | 1.632 (Low)                    | 1.404 (Low)                   | 1.216 (Low)       |        |
| Cu                  | 0.365 (High)                   | 0.188 (Low)                   | 0.446 (High)      |        |

Fig. 1 Composting by using NIPHM bio-inoculum

Kitchen waste cut into bits | Sun drying for 2-3 days | Mixing with NIPHM bioinoculum and poured in drums | Compost under preparation after 20 days | End product after 40 days (Compost) | After ground compost after 50-60 days

Fig. 2 Composting by regular method

Waste vegetable cutting into bits | Bits are ready for mixing with bio inoculum, sanitizer and coco peat | Coco peat bio inoculum and sanitizer | Transferred in to drums | After 30 days wet compost | After 70 days compost ready | Final grounded compost
Fig. 3 Litters after mixing with bio-inoculum covered with polythene sheet

Fig. 4 a. Mixing after 12 days  b. Compost after 60 days

Fig. 5 Growth of isolates on NA medium

Isolate 1  Isolate 2
**Fig. 6** Growth of *Spinacia oleracea* performance in different compost

Growth after FYM compost application  
Growth after vermicompost application  
Growth after vegetable waste compost application  
Growth without compost application

**Fig. 7** Growth of roots of *Spinacia oleracea* in different compost
Growth of *Spinacia oleracea* in different compost

Growth was best in the plot where vegetable waste compost applied followed by vermicompost and FYM compared to control (without compost application). Root length and shoot length were found increased in all the compost planting compared to control. Following figures shows the differences of growth (Fig.6 and 7.)

The time taken for the degradation process to form compost was 50 days. In the case of vegetable wastes, the decomposition process was started after the application of the inoculum which was visualized clearly after 15 days of application by appearance of microbial growth. A sharp decrease in volume and colour change, development of pleasant odour, changes in texture and less water activity was seen. The same observation was seen rapidly in the successive days also. The complete decomposition was clearly observed on the 50th day, which was clearly identified by sharp decrease in volume (3/4th of the volume) complete decolourization, complete absence of water content and complete conversation of finely ground powder which reveal the decomposition of vegetable wastes into fine powder. The temperature of the compost was increasing gradually in the first week and attained maximum on 15th day and started decreasing after wards. After a month it reached to the normal. Composting activity was best at a moisture content of 40-60% by weight. At lower moisture level, microbial activity is very much limited. At higher levels, the process is likely to become anaerobic and foul-smelling.

In conclusion, NIPHM developed the method for composting vegetable waste and leaf litters and also NIPHM developed bio-inoculum which accelerate the decomposition process. The *Spinacia oleracea* was superior to other compost (FYM and Vermicompost). Further more research is required to study the effect of vegetable compost on each component of plant health.

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