# ASSOCIATIVE NITROGEN FIXATION IN SOME GRASSES AND CEREALS GROWING AROUND FAISALABAD AREA

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## ABSTRACT

Nitrogenase activity associated with the roots of five local gresses and cereals i. e. Zea mays L.; Cynodon dactylon, Pers; Sorghum bicolor Pers.; Dactyloctenium aegyptium (L.) Beauv. and Oryza sativa L. was determined using acetylene reduction assay (ARA), for pre-incubated and fresh excised roots. All except O. sativa showed higher nitrogenase activity, both for pre-incubated and fresh roots. Among pre-incubated roots, the unwashed roots showed average nitrogenase activity in the range of 4-107 nmol g-1 D.W.hr-1, washed roots showed in the range of 4-96 nmol g-1D.W. hr-1 and surface sterilized roots activity which was detected in two grasses (Zea mays L. and Cynodon dactylon Pers.), was in the range of 1-9 nmol g-1 D.W. hr-1. For fresh roots, activity was in the range of 2-29 nmol g-1 D.W. hr-1.

#### INTRODUCTION

Nitrogen fixation associated with grasses is not a new concept as Lipman and Taylor (1923) reported this in wheat plant. Recent energy crises and development of sensitive and relatively easier techniques like acetylene reduction assay (Hardy et al., 1968) renewed the interest in this area. Considerable data have been accumulated by using acetylene reduction assay (ARA) by different workers (Dobereiner et al., 1972; Ann Berkum and Day, 1980). Associative nitrogen fixation has also been reviewed by different authors (van Berkum and Bohlool, 1980).

The nitrogen fixing bacteria associated with grass roots have been isolated and identified (Zsfar et al., 1984).

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In this study nitrogenese activity associated with five local grasses and cereals viz. Zea mays; Cynodon dactylon; Sorghum bicolor; Dactyloctenium aegyptium and Oryza sativa has been reported.

## MATERIALS AND METHODS

Collections: The grass root samples were collected from different locations in NIAB fields in the mid-afternoon. The plant roots along with soil cores were dug in a radius of 10 to 20 cm depth (Capone and Taylor, 1980). The samples were immediately brought to laboratory.

Soil Analysis: The rhizosphere soil was analyzed for some of its physicochemical characteristics. The total nitrogen was analyzed by microkjeldahl method (Bremner, 1965). For ammonium and nitrate analysis, 5 of ground soil was shaken with 50 ml of 2NKCI for 1 hr on a mechanical shaker and then filtered. For NH-+.N analysis, 20 ml of the aliquot of the suspension was distilled with MgO (heated at 600°C for 24 hrs) and the distillate was collected in boric acid indicator solution and contents were calculated. For NO<sub>3</sub> —N analysis, 0.20 g finely ground Devorda's alloy (E. Merck) added to the above distillation flask and redistilled. The distillate was collected as described for NH<sub>4</sub> —N. To determine the pH of the soil, a saturated soil paste was prepared in distilled water and pH was determined with pH meter. For EC<sub>6</sub> determinations, the extract from the soil paste was taken and EC<sub>6</sub> was determined using EC meter (WTW- LF-530).

Excised root Assay: The roots were cut into 2-3 cm long segments using sterilized foreceps and scissors. Excised root assay was done for pre-incubated and fresh roots.

- a) Assay with Pre-incubation: The roots were divided into three portions viz. unwashed, washed (washed several times with sterile distilled water) and surface attributed (sterilized with 0.2 HgCl<sub>2</sub> sol, for 30 sec.) roots. The assay was carried out according to the method of Dobereiner and Day (1976).
- b) Assay without Preincubation: Fresh excised roots were taken and subjected to ARA according to the method of van Berkum (1980).

Acetylene Reduction Assay: For estimating the amount of ethylene gas produc-

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ced as a result of acetylene reduction, 100 µl gas mixture was injected into the gas chromatograph (Carlo-Erba Fractorap series 2150) consisting of a 0.75 m x 2 mm stainless ateel column packed with porapak N (80-100 mesh: Water Associate., Inc USA), Nitrogen (Pakistan Oxygen Limited) was used as a carrier gas at a flow rate of 30 ml min-1. The acetylene and ethylene were measured using flame ionization detector (FID). The signal after proper amplification was recorded on Perkin Elmer 56 recorder. The quantity of C<sub>2</sub>H<sub>4</sub> produced was calculated by comparing with standard ethylene in 1% Argon (Linder Technisch Gase, W. Germany). The activity was described in nmol. C<sub>2</sub>H<sub>4</sub> gat D. W. ht-1.

## RESULTS AND DISCUSSION

Soil Analysis: It was known that high concentrations of NH<sub>4</sub> and NO<sub>3</sub> cause reduction in nitrogenase activity (van Berkum and Bohlool, 1980). The soil analysis carried out in these studies is given in Table 1. It showed that total HN, NH<sub>4</sub> N<sub>3</sub> and pH values were almost similar for all grasses but NO<sub>3</sub>- N were high in D. aegyptium followed by Z. mayz and EC<sub>6</sub> values were higher in Z. mayz followed by C. dactylon.

Table 1. The physicochemical characteristics of the soil amples collected from different sites in NIAB fields under various grasses and cereals

| Characteristics            | Z, mays | C. dactylon | 8. bicolor | D. aegyptium | O, sativa |
|----------------------------|---------|-------------|------------|--------------|-----------|
| pH                         | 8.3     | 7.9         | 7.8        | 8.1          | 7.7       |
| ECe (mS. emal)             | 9.56    | 1,47        | 0.94       | 1.24         | 0.73      |
| Total N (mg, g-1)          | 0.063   | 0.062       | 0.052      | 0.053        | 0.032     |
| NH <sub>4</sub> -N (ppm)   | 2.8     | 0.7         | 2.8        | 2.8          | 2,8       |
| NO <sub>3</sub> = -N (ppm) | 5,6     | 0.7         | 2.8        | 7.0          | 1.7       |

To determine the root associated nitrogenase activity both for pre-incubated and fresh roots, a very accurate and sensitive assay called acetylene reduction assay (ARA) of Hardy et al. (1968) was used to get accurate results.

ARA with Pre-incubation: The nitrogenase activity of pre-incubated roots is given in Table 2. The activity was higher i. e., in the range of 4-107 nmol gall D. W. hrall and 4-96 nmol gml D. W. hr. I for rhizosphere (unwashed roots) and rhizoplane (washed roots), respectively but was lower i. e., in the range of 1-9 nmol gall D. W. hrall in histoplane (surface sterilized roots). The histoplane activity was detected only in two grasses i. e., Z. mays and C. dactylon but in others it was not found. It might be due to the experimental error. The histoplane activity was determined due to the reason that there were reports of endorhizosphere nature of certain microhes. The information was taken from light microscopic studies and staining the root tissues with tetrazolium (Patriquin Dobereiner, 1978). In these studies, O. sativa roots showed the lowest activity for pre-incubated roots.

Table 2, Nitrogenase activity associated with preincubated roots of grasses and cereals.

|              | 100000   | 3842     |       |       | Activity | in pmol g                              | · D.W              | , br=t       | 32   |
|--------------|----------|----------|-------|-------|----------|--|--------------------|--------------|--|
| Gras ssystem | L        | Unwashed |       |       | Washed   | ! :::::::::::::::::::::::::::::::::::: | Surface sterilized |              |  |
|              | Freq (%) | Ave      | Range | Freq. | Ave,     | Range                                  | Freq (%)           | Ave,         | Range  |
| Z. maya      | 34       | 18       | 9-27  | 67    | 96       | 6—186                                  | 23                 |              | 414  |
| C. dactylon  | 78       | 107      | 2-211 | 78    | 28       | 1-54                                   | 23                 | 1            | 01   |
| 8. bicolor   | 45       | 20       | 4-36  | 67    | 4        | 1—7                                    | 22                 | -            | _  |
| D. aegyptium | 23       | 13       | 2-25  | 23    | 10       | 2-17                                   | 226                | 2            | 2 <u>.                                    </u> |
| O. sativu    | 23       | 4        | 3-4   | 34    | 7        | 5—9                                    | _                  | 5000<br>5000 | 20-36<br>4 <u>23</u> 866                       |

The present results of pre-incubated roots confirmed those of workers carlier (Neves et al., 1976) who found higher activity with pre-incubated roots of certain grasses.

ARA without Pre-incubation: Earlier, it was considered that pre-incubation period was necessary to detect the nitrogenase activity with fresh excised roots but van Berkum and Sloger (1981) compared the legumes and grasses for nitrogenase activity and they found that both the systems showed C<sub>2</sub> H<sub>2</sub> reduction

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immediately after exposure to  $C_2H_2$ . They concluded that pre-incubation period was not necessary for the detection of nitrogenase activity. Similar results were obtained by De-Polli et al. (1982).

In these experiments, the fresh excised roots were taken for this purpose and subjected to assay according to the method of van Berkum (1980). The results of fresh root assay are given in Table 3. The results of fresh roots assay showed lower values than those of preincubated roots. Similar results were obtained by other workers. The O. sativa roots again showed lower activity than the grasses.

Table 3. Nitrogenase activity associated with fresh roots of grasses and cereals

| Frequency | Activity (nmol g-1)         | D.W. hr=1)                          |  |
|-----------|-----------------------------|-------------------------------------|--|
| (%)       | Average                     | Range                               |  |
| 78        | 28                          | 4-51                                |  |
| 89        | 29                          | 454                                 |  |
| 56        | 4                           | 2 5                                 |  |
| 23        | 13                          | 2-24                                |  |
| 45        | 2                           | 1- 3                                |  |
|           | (%)<br>78<br>89<br>66<br>23 | (%) Average  78 28 89 29 56 4 23 13 |  |

Although the rates of nitrogenase activity associated with roots of grasses are lower as compared to the legumes, yet it carries considerable importance as the grasses occupy about 80% area of the earth hence contributing more nitrogen than legumes. Another important aspect peculiar to grasses is that nearly all grasses have C<sub>4</sub> dicarboxylic pathway of photosynthesis. They leak their photosyntoats in the form of root exudates which is a source of energy for root associated N<sub>2</sub>-fixing microorganisms (Dobereiner et al., 1972). The C<sub>4</sub> grasses utilize their available nitrogen very efficiently in producing dry matter which stimulates the nitrogen fixation in their rhizosphere (Black et al., 1977). Hence the nitrogen fixation with grasses is helpful in food production and to meat the N-demands.

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