

ACTIVITY OF *Azotobacter* AND *Azospirillum* IN THE RHIZOSPHERE OF ONION PLANT

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ABSTRACT

This study was carried out to evaluate the influence of inoculation with nitrogen fixing bacteria (*Azotobacter* and *Azospirillum*) that were isolated from onion rhizosphere. The isolates were applied in field experiment to investigate the effect of inoculation with *Azotobacter* and *Azospirillum*, on Nitrogenase activity. Isolation and enumeration of *Azotobacter* and *Azospirillum* from the soil of onion plants were studied and determination of the enzymatic activities of nitrogen fixing bacteria (nitrogenase) that were isolated from the soil of onion plant during the growing periods at 0, 30, 60, 90, 120 and 150 days after sowing and selected active isolate of *Azotobacter* and *Azospirillum*. The obtained results revealed increases in *Azotobacter* and *Azospirillum* numbers as well as nitrogenase activity in rhizosphere onion regions. Also these activities reflect on onion plant productivity. This study recommended the application of *Azotobacter* and *Azospirillum* as inoculants in onion plant productivity.

Keywords: Isolation *Azotobacter*, *Azospirillum*, Nitrogenase activity.

INTRODUCTION

Onion plants have gotten some interest on seed-inoculated micro-organisms and it was clear that the asymbiotic biological nitrogen fixation was highly correlated to the existence of specific micro-organisms that possess nitrogenase properties such as studied by (Hegazi, *et al.*, 1980, Shoukri, 2006 and Abd el-Halim 2009) that high nitrogenase activity was obtained by the collected samples of *Azospirillum* from plant rhizosphere (Mohy el-Deen 2002 and Narolia, *et al.*, 2006). Nitrogenase activity in onion soil as measured by acetylene reduction technique was found to be high and that indicates positive correlation between *Azotobacter* and higher plants such as onion (Dobereiner *et al.*, 1973 and Chungwoo Kim *et al.*, 2005). Results showed that the isolates of nitrogen fixing bacteria (*Azotobacter* sp. and *Azospirillum* sp.) exhibit a highly activity of nitrogenase enzyme were used as a bacterial inoculants in field experiment, the isolate of *Azotobacter* sp. was 94.09 nmoles C₂H₄/g soil / hr in clayey soil and with the isolate of *Azospirillum* was 10.20 nmoles C₂H₄/g soil / hr in clayey soil. This study aims to find out and enumeration of non symbiotic nitrogen fixing bacteria (*Azotobacter* and *Azospirillum*) from onion rhizosphere, selection of most efficient biological nitrogen fixing bacteria and their influences on onion production at different types of soils.

MATERIALS AND METHODS

Materials:

Onion seeds:

Onion seeds (Giza 6) were kindly supplied by Onion Research Department, Field Crops Research Institute, ARC, and Giza, Egypt.

Experiment This experiment was to study rhizosphere and soil apart of onion plant. This study was carried out in pots (35cm) under green house conditions.

Soil used: The types of the used soils that were collected from the top layer (25 cm depth); a fertile clayey soil and a silty soil were collected from the farm of Malawi Agricultural Research Station. Soils were dried, ground to pass through 2mm sieve, their chemical and physical analyses were presented in Table (1).

Table (1): Physical and Chemical properties of the field soil experiment .

| | | Soil types | | |
|------------------------------------|-------------|-------------------------------|-------|-------|
| | | Clayey | Silty | |
| Physical properties | | pH | 8.90 | 8.20 |
| | | Sand % | 19.90 | 30.50 |
| | | Silt % | 32.60 | 39.00 |
| | | Clay% | 47.50 | 30.50 |
| Chemical properties | Anions (-) | CO ₃ ⁼ | Nil | Nil |
| | | HCO ₃ ⁻ | 2.20 | 1.25 |
| | | Cl ⁻ | 2.00 | 0.90 |
| | | SO ₄ ⁼ | 0.94 | 0.76 |
| | Cations (+) | Ca ++ | 1.30 | 0.50 |
| | | Mg++ | 2.10 | .030 |
| | | Na+ | 1.60 | 1.80 |
| | | K+ | 0.14 | 0.31 |
| | | Organic carbon % | 2.4 | 1.2 |
| Electrical conductivity (mhos/cm) | | 0.56 | 0.28 | |
| C / N ratio | | 3.9 | 2.06 | |
| Total N ₂ % | | 0.19 | 0.11 | |
| Available P % | | 16.8 | 2.11 | |

.A soil analysis was done according to the method described by Jackson, (1973). Analyses were carried out in Soil and Water Institute, ARC, Giza.

Methods

Azotobacter isolates:

Azotobacter isolates used through out the present investigation were isolated from rhizosphere of standing onion plants of 45 days old .

Azospirillum isolates:

Azospirillum isolates were isolated from rhizosphere of the onion plants of 45 days old. One isolate of Azospirillum was isolated from the soil apart, under onion plants.

Isolation and characterization of Azotobacter and Azospirillum from rhizosphere of onion plants

Isolation and purification of the Azotobacter;

Isolation of Azotobacter from the onion rhizosphere was carried out according to the method given by Bilal *et al*, 1990. Ten grams portions from the mixed rhizosphere soil samples were transferred to a sterile 250 ml sampling bottle containing 90ml sterile tap water. The bottles were shaken on a rotary shaker for 10 minutes and further serial dilutions were done. One ml from the appropriate dilution was transferred to 100ml of Stanier basal medium free nitrogen (Stanier, *et al.*; 1963) in flasks 250ml, and incubated at 30°C for 72hrs. from positive tubes, formed pellicle is transferred with sterile loop; streaked surface of basal medium free nitrogen. Incubated at 30°C for 72hrs. The colonies of different morphologies developed on the N-deficient medium were picked up and streaked to obtain a single colony (Abd-El Malek and Ishac, 1968).

Isolation and purification of Azospirillum;-

From root rhizosphere the enrichment cultured technique was adapted using the nitrogen deficient semisolid malate (NFM) recommended by (Dobereiner and Day, 1976) dispensed in test 7ml/tube. Root free soil was homogenized by thoroughly mixing and shaking for 10 minutes first dilution (1:10 w/v) was prepared by transferring 5 ml of roots together with adhering soil in to sampling bottles containing a suitable volume of sterile tap water this method was carried out by Holm and Jensen (1972). Bottles were shaken vigorously for 5-10 min and further serial dilutions were prepared. Isolates of Azospirillum were inoculated in the NFM medium, for enrichment culture for further purification by striking plates to single colony isolation.

Methods of identification of the isolated Azotobacter and Azospirillum strains:-

Nine isolates of Azotobacter and Nine isolates of Azospirillum were tested for cell shape , motility , Gram reaction slime production and pigmentation. All tests were carried out on 24 hrs, old cultures using the nitrogen free medium of (Stanier,*et al*; (1963). The isolates were allowed to grow for 5 days for testing slime production and for 14 days for observation of pigmentation (Bergey's Manual, 2008). The following microbiological tests were followed to identify the isolates:-

Morphology and Gram stain:, Shape, arrangement of the bacterial cells, as well as the Gram reaction were microscopically observed in strain preparations of 24-28hrs.

Motility test:, was observed in fresh preparations 24 hrs by hanging drop technique.

pigmentation:, formation of pigments was reported for 14 days old cultures.

Glucose fermentation:, sterile tubes containing 0.5 % glucose and bromthymol blue (1.0%) as an indicator were inoculated with 0.1 of 24hrs, old cultures. After incubation at 30°C, the change in colour to yellow and the production of gas noted in Durham's tubes.

Starch hydrolysis:, plates of starch agar were streaked with tested isolates and incubated at 30°C for 5 days. Starch hydrolyzing microorganisms were

distinguished by the appearance of clear zones around their growth when the plates were flooded by iodine solution.

Preparation of inocula:

Efficient local strains of *Azotobacter* sp No (5)and *Azospirillum* sp No (1)which had been isolated from the rhizosphere of some onion plants, were used. Heavy cell suspensions of each strain were obtained by growing for 5 days at 29°C, on Ashby and Dobereiner media for *Azotobacter* and *Azospirillum* respectively. Ten ml of *Azotobacter* inoculum-suspension (1.5×10^5 cells/ml of medium) or Ten ml. wer added of *Azospirillum* inoculum-suspension (1.4×10^5 cells/ml of medium) for each plant, as sub-soil biofertilizers, in the rhizosphere area.

Enzymatic activity of nitrogen fixing bacteria:

Nitrogenase activity:

The nitrogenase activity was measured according to (Turner and Gibson, 1980) as follow:

- 1-Sample from the soil of onion plant was collected (25 g) in a 100 ml bottle.
- 2- ml sugar solution were added and well stirred, the solution is composed of 0.5 gram mannitol, 0.5 gram malice acid, 0.5 grams glucose and 0.5 gram sucrose milted in 100 ml distilled water .
- 3-The bottle was closed tightly and removed the air inside it then inject acetylene gas (10 % V) and put a sticker on the injection position.
- 4-The bottle was incubate on 30° C for 24 hours .The acetylene and ethylene were measured in the sample using gas liquid chromatography model HP6890 ,
- 5-The concentration of ethylene in the samples (nmols / C_2H_4 / hr) was then converted to moles by dividing these values by the volume of the molecular weight of gas (22.4 L). The results were presented as n mol / C_2H_4 / ml culture / hr.

RESULTS AND DISCUSSION

Characteristics of the *Azotobacter* and *Azospirillum* isolates:-

The isolation and purification of nine isolates of *Azotobacter* and nine isolates of *Azospirillum* were performed on their specific medium. According to their morphological and physiological characteristics as proposed by Tarrand *et al* (1978) and Bergey's manual (2008). Results recorded in Table (2) illustrated that isolates that developed on Abdel-Malek and Ishac media (1968) have large ovoid to rods shaped, Gram negative, encysted, capsule formation . According to Bergey's manual (2008).The isolates were identified as *Azotobacter spp*. Isolates that grown on NFM medium are vibriod and spiral shape, Gram negative , starch hydrolysis is negative ,motility test is positive , acid from glucose not formed . The data in table (2) showed that the organisms could be indentet as *Azospirillum* According to Bergey's manual (2008) and Chahal, *et al.* (1982) . The most active isolates of *Azotobacter* and *Azospirillum* were used as bacteria inoculum in the field experiment under nitrogen fertilizer levels (0.175 and 350 Kg / fed.)

Table (2); Some morphological and physiological characteristics of Azotobacter and Azospirillum isolates from onion plants .

| Serial number of isolates | | Morphological and physiological characteristics | | | | | |
|---------------------------------------------------|---|-------------------------------------------------|------------|-----------|---------------------------------|------------------------------------------------|-------------------|
| | | Motility | Gram stain | Cell form | Production of acid from glucose | Production of yellow green fluorescent pigment | Starch hydrolysis |
| Picked organisms isolated from N-deficient medium | 1 | - | - | Oval | + | - | - |
| | 2 | - | - | ' | + | - | - |
| | 3 | + | - | ' | + | - | ++ |
| | 4 | + | - | ' | + | + | + |
| | 5 | - | - | ' | + | - | - |
| | 6 | - | - | ' | + | - | + |
| | 7 | + | - | ' | + | + | - |
| | 8 | - | - | ' | + | + | - |
| | 9 | - | - | ' | + | + | + |
| Picked organisms isolated from NFM medium | 1 | + | - | Spiral | + | - | - |
| | 2 | + | - | ' | - | - | - |
| | 3 | + | - | ' | + | - | - |
| | 4 | + | - | ' | - | - | - |
| | 5 | + | - | ' | + | - | - |
| | 6 | + | - | ' | + | - | - |
| | 7 | + | - | ' | + | - | - |
| | 8 | + | - | ' | - | - | - |
| | 9 | + | - | ' | - | - | - |

Data in Table (3) showed that the isolates of nitrogen fixing bacteria (*Azotobacter* and *Azospirillum*) in two soil types exhibit a highly activity of nitrogenase. The activity of the isolate *Azotobacter sp* was 94.09 nmoles C₂H₄ / 1ml culture / hr No 5 in clayey soil , while the isolate *Azotobacter* in *Azospirillum* was 10.20 nmoles C₂H₄ / 1 ml culture /hr No 1 in clayey soil. Such findings confirm those obtained by (Chungwoo Kim *et al.* ;2005 and Abd El-Halim. 2009).

Data presented in Table (4):Showed that the mean counts of *Azotobacter sp.* cells gradually increased in clayey and silty soils with 175 Kg N₂/fed.than soils with 350/kgN₂ /fed In clayey soil ; the highest level of mean count was 8.8 x 10⁴ cells / 1 g dry soil after 60 days from sowing amended with 175 kg N₂/fed compared to control without N₂ fertilizer was 5.1 x 10⁴ cells / 1 g dry soil and to 350 kg N₂/fed was 6.4 x 10⁴ cells / 1 g dry soil after 120 days from sowing compared to control without N₂ fertilizer was 3.8 x 10⁴ cells / 1 g dry soil . In silty soil; the highest level of mean counts with 175 Kg N₂/ fed. was 8.1 x 10⁴ cells / 1 g dry soil after 90 days from sowing compared to control without N₂ fertilizer was 3.4 x 10⁴ cells / 1 g dry soil after 90 days from sowing .The results obtained were in agreement with Kole *et al.* ;1988, Shoukri ; 2006 ,Abd el-Halim. 2009 and Ki-Yoon Kim *et al.*, 2010. *Azospirillum sp.*