



## Effects of *Glomus Mosseae* and *Azospirillum* on The Growth Behavior of Black Gram *Vigna Mungo* (L.) Hepper

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### ABSTRACT

Azospirillum lives in natural soil or in close associations with plants in the rhizosphere. It is helpful to plants and important to farmers because it is able to fix nitrogen (N<sub>2</sub>). It can convert nitrogen gas in the air into nitrogen bound up in amino acids and proteins. Arbuscular Mycorrhizal Fungi (AMF) can increase the capability of the root systems to absorb and translocation of Phosphorus (P) and minor elements through an extensive network of mycelium. AMF are commonly associated with legumes and can increase nutrient uptake of plants growing in high phosphate fixing soils. The present study the efforts of single and dual inoculation of *Glomus mosseae* and *Azospirillum* on black gram *Vigna mungo* (L.) Hepper. The AMF *Glomus mosseae* + *Azospirillum* inoculation plants resulted in production of highest biomass such as plant, shoot length, root length, number of leaves, number of root nodules, fresh and dry weight of black gram plants and the biochemical content such as total chlorophyll, total sugar and protein. The nutrient contents such (macro and micro nutrients). The inoculation of *Glomus mosseae* + *Azospirillum* showed an enhanced among the parameters when compared to control seedlings.

### KEYWORDS

*Glomus mosseae*, *Azospirillum*, Phosphorus, Nitrogen, AMF, Morphological and Biochemical parameters.

### Introduction

Biofertilizer is defined as a substance which contains living microorganisms and is known to help with expansion of the root system and better seed germination. The microorganisms containing biofertilizers can be the tools we could change apply of chemical fertilizers. Biofertilizers are products containing living cells of different types of microorganism, which have an ability to convert nutritionally important elements to available form through biological processes. In recent years, biofertilizers have emerged as an important component of the integrated nutrient supply system and hold a great promise to improve crop yield through environmentally better nutrient supplies (Marianna *et al.*, 2005). There is a great interest in establishing novel associations between higher plants and various N<sub>2</sub>-fixing microorganisms and Phosphorus solubilizing fungi (Al-Khiat, 2006).

*Azospirillum* are gram-negative to gram-variable, curved-rod shape, motile, oxidase positive and exhibit acetylene-reduction activity (ARA) under micro-aerophilic conditions. *Azospirillum* spp. has been identified mainly as rhizosphere bacteria and its colonization of the rhizosphere has been studied extensively along with reporter gene fusion (Pereg-Gerk *et al.* 2000). *Azospirillum* spp. is a most studied plant growth promoting bacteria because of its ability to colonize in roots of different plant species and most of the plants are of agricultural importance (Cessán *et al.*, 2008). These bacteria use atmospheric nitrogen for the synthesis of the cellular proteins. Cellular protein is mineralized after the death of the cell, thus contributing to the availability of nitrogen for wild plants and crops (Agronet 2012).

AMF, are particularly ubiquitous in soil and create symbiotic associations with most terrestrial plants including agricultural crops, cereals, vegetables, and horticultural plants. In agriculture, several factors such as host crop dependency to mycorrhizal colonization, tillage system, fertilizer application, and the potential of AMF inoculum, affect plant response and plant benefits from mycorrhizae. (Swift, 1998) Interest in AMF propagation for sustainable agriculture is increasing due to

its role in the promotion of plant health, and improvements in soil fertility and soil aggregate stability. These fungi can be utilized effectively for increasing yields while minimizing use of pesticides and inorganic fertilizers. (Bethlenfalvai, 1992).

N<sub>2</sub> is an integral component of many compounds including chlorophyll and enzymes essential for plant growth processes. It is essential component of amino acid and related proteins. N<sub>2</sub> is essential for carbohydrate use within plants and stimulates root growth and development as well as the uptake of other nutrients. This element encourages above ground vegetative growth and gives a deep green colour to the. It is recognized that nitrogen is one of the key elements of soil fertility leaves (Brady and Weil 2002). Blackgram being a leguminous crop is capable to fix atmospheric N<sub>2</sub> through symbiosis.

Phosphorus (P) is vital to plant growth and is found in every living plant cell. It is involved in several key plant functions, including energy transfer, photosynthesis, transformation of sugars and starches, nutrient movement within the plant and transfer of genetic characteristics from one generation to the next. Promotes early root formation and growth, Improves quality of fruits, vegetables, and grains, Increases water-use efficiency (Ahmad *et al.*, 2009)

Black gram is the third important pulse crop in India. It is annual pulse crop and native to central Asia. It is also extensively grown in West Indies, Japan and other tropics and subtropical countries. The seeds are highly nutritious containing higher amount of protein (24-26 %) and rich in potassium, phosphorus and calcium with good amount of sodium. It is also reported to be rich in vitamin A, B<sub>1</sub>, B<sub>3</sub> mineral and vitamins. It has some medicinal properties, like curing diabetes, sexual dysfunction, nervous disorder, hair disorders, digestive system disorders and rheumatic afflictions (Anonymous, 2010)..

**Materials and Methods**  
Seed

The seeds black gram *Vigna mungo* (L.) Hepper varieties (IPU-941) were obtained from Local Agro seeds centre Dharmapuri Taluk and Dharmapuri district, Tamil Nadu, India. The seeds uniform size, colour and weights were chosen for experimental purpose. Seeds were surface sterilized with 0.1 percent mercuric chloride solution and washed thoroughly with tap water and then with distilled water.

#### AMF and *Azospirillum* Collection and Inoculation;

The AMF and *Azospirillum* were collected at District Forest Office (Modern Nursery Division) the AMF species employed was *Glomus mosseae*. The inoculum used consisted of Vermiculite soil containing spores (800 to 1000 / 200g dry soil), hyphal fragments and fine roots of maize infected with *Glomus mosseae*. *Azospirillum* was applied @ 1 kg ha<sup>-1</sup> by mixing with seed.

#### Pot culture experiments

Twenty five seeds of black gram were sown in each earthen pots separately were filled with 5 kg/pot of sandy and Red loam soil. The experiment was arranged as a randomized black design with five replicates for each treatment. The following replicates were carried out with the following treatments. T<sub>1</sub>- Control, T<sub>2</sub> - AMF, (*G.mosseae*) T<sub>3</sub> - *Azospirillum* and T<sub>4</sub> - *G.mosseae* + *Azospirillum*

Black gram plants were collected at 50<sup>th</sup> days and their shoot and root length was measured and recorded. The roots were dipped in water to remove adhering soil particles and washed with tap water and distilled water the number of root nodules were estimated. Dry weights of the plants were determined after drying at even 80°C of a constant weight. The chlorophyll content was assayed according to (Li, 2000) the extraction was made from a 100 mg fresh sample in 25 mL acetone (80%) in the dark at the room temperature and measured at 470, 646 and 663 nm with a UV spectrophotometers. The sugar and protein content of plant leaf materials were estimated according to (Naguib 1963) and (Bradford, 1976). Nitrogen (N) content was extracted from sulfuric acid using the semi micro kjeldhal method (Jockson *et al.*, 1973). Phosphorus (P) was extracted by nitric acid and perchloric acid digestion and measured using the vanadoso-molybdophosphoric colorimetric method (Jockson, 1967). Potassium (K) was assayed using a flame spectrophotometer (Allen *et al.*, 1984) calcium (Yoshida *et al.*, 1972), magnesium (Jackson, 1958), Zinc, copper, Sulphur, iron and manganese (Piper, 1966)

#### Results and Discussion

Biofertilizers will be the best solution to replace chemical fertilizers. Biofertilizers are the carrier-based arrangements containing mostly valuable strains of microorganisms in adequate number, which are useful for N<sub>2</sub> fixation. Amongst the nutrients, nitrogen is the only nutrient, which play major role in synthesis of chlorophyll, amino acids and protein building blocks (Bloemberg 2000).

The pot experiment was carried out to inoculation with *G. mosseae* and *Azospirillum* on morphological and biochemical contents were presented in Table 1. The experimental results, maximum values of root length (29.8cm), shoot length (42.7cm), No of leaves (47.4), total leaf area (37.2), number of root nodules (81.5), fresh weight (48.52 g), Plant dry weight (23.64g) total chlorophyll (4.6 mg.fr.wt), total sugars (7.6) and protein (18.28 mg.fr.wt) was recorded in AMF + *Azospirillum* application at 50 DAS of black gram plant. The minimum root length (22.5cm), shoot length (34.2cm), No of leaves (38.2), total leaf area (29.7), number of root nodules (51.2), fresh weight (40.48 g), Plant dry weight (18.43g) total chlorophyll (3.22 mg.fr.wt), total sugars (3.39) and protein (9.5 mg.fr.wt) was recorded in control treatments at 50 DAS of black gram plants.

The data indicated that the dependence of black gram plant growth on inoculation with *G. mosseae* + *Azospirillum* the increased the root and shoot length on all sampling days. The increase in plant height might be attributed to the N<sub>2</sub>-fix-

ation by *Azospirillum* which inturn make the essential nutrients available to the plant growth and development. These substances have also been reported to increase the activity of cell division and cell elongation ultimately leading to an increased plant height. Similar results have also been reported from Fallik and Okon (1996) in *Setaria italica*, cauliflower as Jawar moti (Kalyani *et al.*, 1996). Arbuscular Mycorrhizal Fungi (AMF) symbioses have been shown to benefit growth of many field crops in large part due to the extensive hyphal network development in soil, more efficient exploitation of nutrients, and enhanced plant uptake (Smith and Read, 1997). The AMF infection is recognized to augment plant growth by increasing nutrient uptake. The higher height increment registered with dual inoculated plants could be as a result of enhanced both organic and inorganic nutrient absorption and greater rates of photosynthesis which obviously could have given to an increase in plant growth (Cooper, 1984).

The *G. mosseae* + *Azospirillum* inoculated plants were increased in the fresh and dry weights of black gram plants. This result was in correspond to (Kirshna and Bagyaraj, 1984) who reported that inoculation with the mycorrhizal fungus *Glomus fasciculatum* enhanced peanut growth and increased its dry matter more than 2-fold compared with control. AMF inoculated plants have the fungal hypha increase root surface area, resulting in exploring higher volume of soil and overcoming the water and nutrient depletion zones around the roots leading to increased water and nutrient content (Clark and Zeto, 2002). Inoculation of plants with *Azospirillum* could result in significant changes in various growth parameters, such as increase in plant biomass, nutrient uptake, plant height, number of leaves, leaf area and seedlings length (Bashan *et al.*, 2004).

The pigment content of total chlorophylls and biochemical content such as total sugars and protein content was significantly increased the dual inoculation of *G.mosseae* + *Azospirillum*. The effect of N<sub>2</sub> fixing and phosphate solubilizer microorganisms exhibited significant differences in total chlorophyll content. Among the treatments, the total chlorophyll content was significantly higher in the treatment AMF + *Azospirillum* at 50<sup>th</sup> days of black gram plants. This could be due to the beneficial effect of dual inoculants which helped the plants to get more nitrogen which is the component of chlorophyll molecule. The delayed leaf senescence has also been attributed to higher total chlorophyll content (Hyakumachi and Kubota, 2004b).and Free-living nitrogen-fixing bacteria eg *Azotobacter chroococcum* and *Azospirillum lipoferum*, were found to have not only the ability to fix nitrogen but also the ability to release phytohormones similar to Gibberellic Acid (GA) and Indole Acetic Acid (IAA), which could stimulate plant growth, and photosynthesis (Fayez *et al.*, 1985).

Sugar is an important energy constituent needed for all living organisms. The concentrations of soluble sugar indicate the physiological activity of plant organisms. The total sugar content was higher in combined inoculation of microorganisms similarly the control plants have lowest sugar content. Increased accumulations of sugars in black gram plant is due to *G. mosseae* + *Azospirillum* application is in conformity with the earlier studies in different micro organisms on crop plants such as maize (Tejeda and Gonzalez, 2006), *Albizia lebbek* (Kumudha and Gomathinayagam, 2007), Suresh and Bagyaraj (1984) reported that AM inoculation increased the quantities of sugars and amino acids in plant tissue.

Protein is one of the reserve food materials which are utilized for the growth of seedlings. The highest protein content of black gram plant was recorded in the treatment of *G. mosseae* + *Azospirillum* application when compared to control plants. The significant increasing in protein content is due to the increase the percentage of 'N' and 'P' in plants (Vazquez *et al.*, 2000). It was found that organic acid of soils increased the plant uptake of P form a water soluble and also the release of organic acids both sequester cations and acidity. The microenvironment near the roots is through to be major

mechanisms of 'P' uptake as well as Mn, Fe and Zn by plants and AM fungi. Increased levels of protein in the inoculated plants could be attributed to either the presence of fungal proteins stimulation of protein synthesis in the host plant (Krishna and Bagyaraj, 1984). It also observed by Arines *et al.* (1993) in red clover. Nanthakumar and Veeraraghavathatham (2000) in brinjal and Govindarajan and Thangaraju (2001) in chilli.

The nutrient uptake was more in dual inoculation of *G. mosseae* + *Azospirillum* application of black gram plants (Table.2). AMF symbioses have been shown to benefit growth of many field crops in large part due to the extensive hyphal network development in soil, more efficient exploitation of nutrients, and enhanced plant uptake. They improve nutrient uptake, especially P, and also uptake of micronutrients such as zinc or copper; they stimulate the production of growth substances and may reduce stresses, diseases or pest attack (Smith and Read, 1997). The role of AMF on nutrient uptake (N, P and

microelements), on the growth of AM crops, as well as on possible mechanisms of nutrient uptake, have been widely studied, as Al-Karaki (2006), Cardoso and Kuyper (2006), and Cavagnaro (2008). It is now generally recognized that AMF enhance the uptake of nitrogen (N) and of relatively immobile soil nutrients such as phosphorus (P), sulfur (S), copper (Cu), zinc (Zn), and boron (B) Martin *et al.* (2007).

AMF formation is known to enhance nodulation and N<sub>2</sub> fixation by legumes (Andre *et al.*, 2005). Mycorrhizal and other symbioses often act synergistically on infection rate, mineral nutrition and plant growth (Amora-Lazcano *et al.*, 1998). The positive fungal effect on plant P uptake is beneficial for the functioning of the nitrogenase enzyme of the microbial symbiont leading to a higher N<sub>2</sub> fixation and, consequently to a better root growth and mycorrhizal development (Johansson *et al.*, 2004). It concluded of my research article for better understanding of soil system would probably lead to a better management of AMF+ *Azospirillum*, contribution to soil fertili-

ty and, more sustainable agriculture, even in high yielding of black gram productions.

Treat-ments	SL (cm)	RL (cm)	No. of L	LA (cm <sup>2</sup> )	No. of RN	FW (mg/g fr. wt.)	DW (mg/g dr. wt.)	T.Chl (mg/g fr. wt.)	T.Sug-ar(mg/g fr. wt.)	Protein (mg/g fr. wt.)
T1	34.2 ± 1.71	22.5 ± 1.125	38.2 ± 1.91	29.7 ± 1.485	51.2 ± 2.56	40.48 ± 2.024	18.43 ± 0.97	3.22 ± 0.161	3.39 ± 0.17	9.5 ± 0.47
T2	37.7 ± 1.85	24.8 ± 1.24	41.5 ± 2.075	33.5 ± 1.675	68.5 ± 3.425	42.1 ± 2.105	19.82 ± 0.091	3.89 ± 0.1945	5.98 ± 0.30	11.21 ± 0.57
T3	38.8 ± 1.89	26.3 ± 1.315	44.2 ± 2.21	35 ± 1.75	77.6 ± 3.88	43.85 ± 2.1925	21.5 ± 1.175	4.08 ± 0.204	6.31 ± 0.32	14.79 ± 0.74
T4	42.7 ± 1.91	29.8 ± 1.49	47.4 ± 2.37	37.2 ± 1.86	81.5 ± 4.075	48.52 ± 2.426	23.64 ± 1.282	4.6 ± 0.23	7.6 ± 0.38	18.28 ± 0.92

**Table.1 Effects of *Glomus mosseae* and *Azospirillum* on the growth and biochemical content of Black gram *Vigna mungo* (L.) Hepper on 50<sup>th</sup> days of plants.**

± Standard deviation

T1 – Control, T2 = *Azospirillum*, T3 = AMF T4 = AMF + *Azospirillum*.

S. L= Shoot length R. L= Root Length No. of L= Number of Leaves LA= Leaf Area FW= Fresh wight No. of. RN= Number of root nodules DW= Dry weight T. Ch l= Total Chlorophyll T.sugar = Total Sugar.

**Table.2 Effects of *Glomus mosseae* and *Azospirillum* on the Nutrient content (mg/g dr. wt.) of Black gram *Vigna mungo* (L.) Hepper on 50<sup>th</sup> days of plants.**

Treatments	N	P	K	Ca	Mg	S	Zn	Cu	Mn	Fe
T1	89.6 ± 4.45	5.5 ± 0.275	0.58 ± 0.03	1.85 ± 0.092	0.89 ± 0.044	1.3 ± 0.11	1.66 ± 0.08	0.14 ± 0.007	0.45 ± 0.023	1.15 ± 0.057
T2	92.7 ± 4.63	6.2 ± 0.31	1.38 ± 0.07	1.54 ± 0.077	1.08 ± 0.054	1.89 ± 0.19	2.18 ± 0.11	0.15 ± 0.0075	1.01 ± 0.051	2.67 ± 0.1335
T3	96.8 ± 4.84	7.9 ± 0.395	2.48 ± 0.2	1.63 ± 0.081	1.74 ± 0.09	2.05 ± 0.22	2.95 ± 0.15	0.24 ± 0.012	1.91 ± 0.096	2.98 ± 0.149
T4	99.2 ± 4.96	8.5 ± 0.425	3.63 ± 0.181	2.48 ± 0.124	2.03 ± 0.102	2.52 ± 0.26	3.68 ± 0.18	0.32 ± 0.016	2.68 ± 0.134	3.79 ± 0.190

± Standard deviation

T1 – Control, T2 = *Azospirillum*, T3 = AMF T4 = AMF + *Azospirillum*.

## REFERENCE

1. Agronet. 2012. Azotobacter. Obtained from: [http://www.indiaagronet.com/indiaagronet/Manuers\\_fertilizers/Manure\\_Fe\\_rt.htm](http://www.indiaagronet.com/indiaagronet/Manuers_fertilizers/Manure_Fe_rt.htm).
2. Ali Khan, A, G. Jilani, M. S. Akhtar, S. M. Saqlan Naqvi, M. Rasheed. 2009. Phosphorus Solubilizing Bacteria: Occurrence, Mechanisms and their Role in Crop Production. *J of Agri. Bio. sci.* 11:48-58.
3. Al-Karaki, G.N., 2006. Nursery inoculation of tomato with arbuscular mycorrhizal fungi and subsequent performance under irrigation with saline water. *Sci. Hort.* 109: 1–7
4. Al-Khiat S and H. Ali, 2006. Effect of cyanobacteria as a soil conditioner and biofertilizer on growth and some biochemical characteristics of tomato (*Lycopersicon esculentum* L.) Seedlings. Thesis Submitted in partial fulfillment of the requirements of the Degree of Master of Science (M. Sc.) Microbiology (Algae), King Saud University. Special Publication, 6, p: 1-4
5. Allen, S.F., H.F. Grimshaw and A.B. Rowland. 1984. Chemical analysis. In : Methods in plant ecology, pp. 185-344. Eds. Moore, P.D. and S.B. Chapman Blackwell, Oxford.
6. Amora-Lazcano, E., M.M. Vazquez and R. Azcon, 1998. Response of nitrogen-transforming microorganisms to arbuscular mycorrhizal fungi. *Biol. Fert. Soils* 27: 65–70.
7. Andre, S, A. Galiana, N. Le, C. Roux, Y. Prin, M. Neyra, and R. Duponnois, 2005. Ectomycorrhizal symbiosis enhanced the efficiency of inoculation with two Bradyrhizobium strains and Acacia holosericea growth. *Mycorrhiza* 15: 357–364.
8. Anonymous, 2010. <http://wik.onse.com>
9. Arines, J, A. Vilarino and M. Sainz 1990. Effect of vesicular-arbuscular mycorrhizal fungi on Mn uptake by red clover. *Agri. Ecosys. Environ.* 29: 1–4.
10. Bashan, Y and G. Holguin, 1998. Proposal for the division of plant growth-promoting rhizobacteria into two classifications: biocontrol-PGPB (plant growth-promoting bacteria) and PGPB, *Soil Biol. Biochem.* 30: 1225-

- 1228.
11. Bethlenfalvay, G.J., 1992. Mycorrhizae and crop productivity. In: Bethlenfalvay, G.J., and Linderman, R.G. (eds.), Mycorrhizae in Sustainable Agriculture. Madison, WI: ASA Special Publication No. 54, pp. 1–27.
  12. Bloomberg GV, A.H.M. Wijffjes, G.E.M. Lamers, N. Stuurman and B.J.J. Lugtenberg 2000. Simultaneous imaging of fluorescent proteins in the rhizosphere: new perspective for studying microbial communities. *Mol. Plant Mic. Int.*, 13: 1170-1176
  13. Bradford, M.M., 1976. A rapid and sensitive method for the quantifications of microgram quantities of protein utilizing. The principle of protein-dye binding. *Anal. Biochem.*, 72: 248-254.
  14. Brady, N.C., and R.R.Weil, 2002. The nature and properties of soils, Prentice Hall, New Jersey, pp. 960.
  15. Cardoso, I.M., and T.W. Kuyper 2006. Mycorrhizas and tropical soil fertility. *Agri. Ecosys. Environ.* 116: 72–84.
  16. Cassan, F., V. Sgroj, D. Perrig, O. Masciarelli and V. Luna. 2008. Phytohormone production by *Azospirillum* spp. physiological and technological aspects of plant growth promotion. In *Azospirillum* spp. *cell phy, plant inter. and agro. res. in Arg*, 61-86.
  17. Cavagnaro, T.R. 2008. The role of arbuscular mycorrhizas in improving plant zinc nutrition under low soil zinc concentrations: A review. *Plant Soil* 304: 315–325.
  18. Clark, R.B. and S.K. Zeto, 2002. Arbuscular mycorrhiza: Mineral nutrient and water acquisition. In : Sharma, A.K., B.N. Johri (Eds.) Arbuscular mycorrhiza, Interactions in Plants, Rhizosphere and soils. *Oxford and IBH Publishing Co. Pvt. Ltd., New Delhi*, pp. 159-188.
  19. Cooper, K.M., 1984. Physiological of VA mycorrhizal association in: VA mycorrhiza (Ed. B.Y.C.L Powell and D.J. Bagyaraj) pp. 155-186. *CRC Press, Inc., Boca Raton. Florida*.
  20. Fallik, E. and Y. Okon (1996): Inoculants of *Azospirillum brasilense*: biomass production, Survival and growth promotion of *Setaria italica* and *Zeamays*. *Soil Biol. Bio chem.* 28, 123–126.
  21. Fayez M., N.F. Emam and H.E. Makkboul 1985. The possible use of nitrogen fixing *Azospirillum* as biofertilizer for wheat plants. *Egypt. J. Microbiol.*, 20(2), 199-206.
  22. Govindarajan, K and M. Thangaraju. 2001. *Azospirillum* - a potential inoculant for horticulture crops. *South Indian Hort.* 49,223- 235.
  23. Hyakumachi, M and M. Kubota 2004. Fungi as plant growth promoter and disease suppressor. In: Arora, D.K. (ed.), Fungal Biotechnology in Agricultural, Food, and *Environmental Applications*. *New York: Marcel Dekker*, pp. 101–110.
  24. Jackson, M.L., 1958. Soil chemical analysis, Prentice Hall of India Private Limited, New Delhi, pp. 22-31.
  25. Jackson, M.L., 1967. Soil chemical analysis. Prentice-Hall, New Delhi, India.
  26. Jackson, M.N.F., R.H. Miller and R.F. Forklin, 1973. The influence of VAM on uptake of 90 Sr from soil by soybeans. *Soil Biol. Biochem.*, 5: 205-212.
  27. Johansson, J.F., L.R. Paul, and R.D. Finlay, 2004. Microbial interactions in the mycorrhizosphere and their significance for sustainable agriculture. *FEMS Microbiol. Ecol.* 48: 1–13.
  28. Kalyani, D.P., C. Ravishankar, and P.D. Manohar, 1996. Studies on the effect of nitrogen and *Azospirillum* on growth and yield of cauliflower. *South India Horti*, 44(5-6): 147-149.
  29. Krishna, K.R. and D.J. Bagyaraj, 1984. Growth and nutrient uptake of peanut inoculated with the mycorrhizal fungus *Glomus fasciculatum* compared with non-inoculated. *Plant Soil*, 77: 405-408.
  30. Kumudha, P. and M. Gomathinayagam, 2007. Studies on the effect of biofertilizers on germination of *Albizia lebbek* (L.) Benth. seeds. *Adv. Plant Sci.*, 20: 417-421.
  31. Li, H.S., 2000. Principles and techniques of plant physiological biochemical experiment. Beijing. *Higher Education Press*.
  32. Marianna, M, S.V E. Gajdos, N. Bakonyi, B. Toth, and L. Levai 2005. The possible role of biofertilizers in agriculture. *Ratarstvo*, 585-588.
  33. Martin, F, Perotto, S and P. Bonfante, 2007, Mycorrhizal fungi: A fungal community at the interface between soil and roots, pp. 201–236. In R. Pinton, Z. Varanini, and P. Nannipieri (Eds.), *The rhizosphere: Biochemistry and organic substances at the soil-plant interface*. Marcel Dekker, New York.
  34. Naguib, M.I., 1963. Colorimetric estimation of plant polysaccharides *Zucker*, 16: 15-18.
  35. Nanthakumar, S and D. Veeraghavathatham. 2000. Effect of integrated nutrient management on growth parameters and yield of brinjal (*Solanum melongena* L.) cv. PLR-1. *South Indian Hort.* 48, 31 35.
  36. Pereg-Gerk L, K. Gilchrist and I.R. Kennedy 2000. Mutants with enhanced nitrogenase activity in hydroponic *Azospirillum brasilense*-wheat associations. *Appl Env. Microbio.* 66, 2175–2184.
  37. Piper, C., 1966. Soil and plant analysis. Asian Hans Publishers, Bombay, pp.11-36.
  38. Smith, S.E. and D.J. Read, 1997. Mycorrhizal symbiosis academic press, San Diego, pp. 605.
  39. Suresh, C. K., and D. J. Bagyaraj 1984. Interaction between vesicular-arbuscular mycorrhizae and a root-knot nematode and its effect on growth and chemical composition on tomato. *Nematol. Medit.* 12: 31–39.
  40. Swift, M.J., 1998. Toward the second paradigm: integrated biological management of soil. Paper presented for the FERTBIO Conference, Brazil.
  41. Tejada, M. and J. Gonzalez, 2006. Effect of foliar application of beet vinase on maize yield. *Biol. Agric. Hort.*, 24: 197-214.
  42. Vazquez, M.M., S. Cesar, R. Azcon, and J.M. Barea. 2000 Interactions between arbuscular mycorrhizal fungi and other microbial inoculants (*Azospirillum*, *Pseudomonas*, *Trichoderma*) and their effects on microbial population and enzyme activities in the rhizosphere of maize plants. *Appl. Soil Ecol.* 15: 261–272.
  43. Yoshida, S., D. Fordo, J. Cork and K. Gomez, 1972. Laboratory manual for physiological studies of rice 3<sup>rd</sup> edn., The International Rice Research Institute, Philippines, pp.11-23.