

Impact of Fertigation on Soil Microbial Community and Enzyme Activities Cropped with Maize (cultivar. COMH 1) under Precision Farming System

M. Jeya Bharathi*, D. Balachandar, R. Narayanan and K. Kumar

Department of Agricultural Microbiology Tamil Nadu Agricultural University, Coimbatore -641 003

Precision farming is an integrated plant nutrient management system attempt to deliver the inputs actual crop needs without disturbing the entire farming area. The fertilizers in liquid state will be delivered through laterals directly to the root zone at high concentrations. Soil microorganisms are potential resources for soil productivity and sustainability, intended to be influenced by various anthropogenic-and farming activities. Hence in present study, we have investigated that the effect of different dose of liquid fertilizers on changes in soil microbial populations and enzyme activities of maize in comparison with conventional practice of fertilization. The results clearly revealed that all the fertilization reduced the microbial population and enzyme activities of maize rhizosphere and among the fertigations, 75 per cent of recommended NPK in the form of liquid fertilizer had a less reduction in the population of *Azospirillum*, P solubilising bacteria and total diazotrophs, soil respiration and enzyme activities. Hence it is evident from the study that the use of more organic manures and microbial inoculants along with inorganic chemicals fertilizers, as integrated approach is needed for precision farming.

Key words: Azospirillum; Fertigation; Maize; Precision farming; Soil enzymes

Precision farming or precision agriculture aims at increasing productivity, decreasing production costs and minimizing the environmental impact of farming (Maheswari et al., 2008). It is an integrated crop management system that attempts to match the kind and amount of inputs with the actual crop needs for small area within a farm field (Ahlwaalia et al., 1993). It is based on soil, weather and crop requirement to maximize sustainable productivity, quality and profitability. Agricultural sustainability requires optimal use and management of soil fertility, soil physical properties, both of which relay on soil biological processes (Saxena and Tilak, 1998) which has been achieved through precision farming. Soil microorganisms are responsible for breakdown of organic matter, help releases of nutrients and their availability for other organisms (Jenkinson, 1988). Further microbial activity and biomass dynamics help to regulate long term soil properties such as net fluxes and amount of soil carbon and nutrients (Bauhus and Barthel, 1995), hence it is served as an important reservoir of plant nutrient such as N and P. The soil microbial biomass is often regarded as an early indicator of changes which may occur in the long term with regard to soil fertility and agro ecosystem properties (Powlson et al., 1987). Its size and activity directly related to the amount and quality of carbon and other nutrients

available from plant residues, organic amendments and root exudates. Soil enzymes are derived primarily from soil fungi, bacteria, plant roots, microbial cells, plant and animal residues, etc. (Brown, 1973) and play a significant role in mediating biochemical transformations involving organic residue decomposition and nutrient cycling in soil (McLatchey and Reddy, 1998).

Management practices which are associated with intensification of agriculture (Giller *et al.*, 1997) are well known to alter soil microbial biomass and activity and this has been the topic of considerable research effort over the past two decades. Thus microbial biomass responds to alteration in tillage practices (Lynch and Panting, 1980) addition of fertilizers (Fauci and Dick, 1994) and pesticides (Wardle and Parkinson, 1991) and manipulation of organic residues (Franzluebbers *et al.*, 1995) and cover species (Beese *et al.*, 1994).

It has been shown that microbial activity and biomass are higher in fields with organic amendments than field with conventional fertilizers (Gunapala and Scow, 1998). The amount of soil nitrogen in fields under conventional production system has been negatively correlated with soil microbial components, where as soil nitrogen in fields under organic production was positively

^{*}Corresponding author email: jbharathi86@gmail.com

correlated with soil microbial components (Gunapala and Scow, 1998). Inorganic N fertilization can have significant effect on soil microorganisms and enzymes through higher plant yield and thus, crop residues and through its impact on soil pH depending on the amount and type of fertilizers (Tabatabai et al., 1992). Nitrogen fertilizers application may lead to lowering the soil pH (Aref and Wander, 1998). Most N containing inorganic fertilizers, unless specially treated tends to acidify soil. This is mainly due to the fact that most fertilizer supply NH4⁺ or result in its production. Upon oxidation NH4⁺ can release H⁺ ions which are potential source of acidification (Magdoff et al., 1997).

Limited field studies have been conducted to determine the impact of soil amendments on microbial communities in actual organic and conventional production systems in the fields (Gunapala and Scow, 1998). As precision farming aims to feed the liquid form of mineral nutrients directly in the rhizosphere region of crop plants, certainly they may cause flux in the native populations of functional microorganisms and enzymes in rhizosphere. With this hypothesis, the present investigation was carried out to study the effect of fertigation on soil microbial community and enzyme activity.

Materials and Methods

Field experiment

A field experiment was conducted to study the impact of liquid fertilizers on soil microbial community and enzyme activity with maize crop (Cultivar COMH 1) under precision farming system at Eastern Block of Tamil Nadu Agricultural University, Coimbatore during Kharif season, 2007. The physic chemical and microbiological properties of the experimental field soil were presented as Table 1. The experiments were laid out in a randomized block design with five treatments and three replications with a plot size of 90 sq.m each. The treatments includes T₁-Uninoculated and unfertilized control, T2-75% RDF (Recommended dose of fertilizer) of NPK through drip, T₃-100% RDF of NPK through drip, T₄- 150% RDF of NPK through drip and T5-100% RDF of NPK through conventional method of application. Fertilizers were applied at weekly intervals in the form of urea, multi K and polyfeed as per the schedule recommended for precision farming system through fertigation tank. In the case of conventional method, fertilizers were applied by broadcasting.

Soil microbial populations

The rhizosphere soil samples at weekly interval were collected at five different plants of each plot and pooled. The stones and stubbles removed soil samples packed in air tight polybags were stored at 4°C for further analysis. The population of *Azospirillum*

was enumerated by MPN count as described by Okon *et al.* (1977). The total diazotrophic bacterial count was done by following the procedure described by Watanabe and Barraquio (1980). The population of P solubilising bacteria was enumerated on hydroxyl apatite medium (Sperber, 1958).

Soil respiration rate

The rhizosphere soils collected at vegetative, flowering and harvest stage of the crop were quantified for soil respiration rate by assessing the amount of CO₂ evolved by alkali trap method (Pramer & Schmidt, 1956) and expressed as mg of CO₂ evolved per g soil per h.

Soil enzymes

Acid phosphatase (E.C. 3.1.3.2) and alkaline phosphatase (EC 3.1.3.1) were determined according to Tabatabai (1982) and reported as μ g p-nitrophenol per gram dry weight of soil. Dehydrogenase activity (E.C.1.2.1.3) was determined by monitoring the rate of production of triphenylformazon (TPF) using the method of Klein *et al.* (1971) and expressed as μ g of TPF per g soil per day. The soil nitrogenase (E.C. 3.2.1.2) was measured by acetylene reduction assay as described by Bergerson (1980) and expressed as μ mol of ethylene produced per g soil per h. Soil urease (E.C. 3.5.1.5) was determined according to Tabatabai and Bremner (1972) and reported as μ g of NH₄ -N released per g soil per day.

Results and Discussion

The fertigation effects in precision farming of maize on soil microbial and enzyme activities in rhizosphere was investigated in present study and the results clearly revealed that significant difference

Table	1.	Physiochemical	properties	of	experi
menta	l pl	ot soil			

Properties	Mea	an ± SE
pH	6.36	± 0.09
EC (ds/m)	1.6	6 ± 0.15
Organic carbon (%)	0.26	± 0.01
Available N (kg/ha)	310	± 2.88
Available P (kg/ha)	44	± 0.57
Available K (kg/ha)	935.3	± 1.45
Total bacteria (cfu x 10 ⁵ / g dry weight of soil) ^a	19	± 1.15
Fungi (cfu x 10 ³ / g dry weight of soil) ^b	4	± 0.16
Diazotrophs (cfu x 10 ⁴ / g dry weight of soil) ^c	46	± 1.15

Values are mean ± SE of three replications

^aTotal bacteria were enumerated by serial dilution plating method on soil extract agar medium (James 1958).
^bTotal culturable fungi were enumerated by serial dilution plating method as described by Parkinson *et al.* (1971).

^cTotal diazotrophs were enumerated by the procedure as described by Rennie (1981).

was noticed between biofertigation and conventional fertilizer application.

Effect of fertigation on rhizosphere microbial population

In general, application of chemical fertilizers stimulated the growth and multiplication of

		CO ₂ evolution (mg CO ₂ /g/h)			
	Ireatment	Vegetative growth stage	Flowering stage	Harvestingstage	
T_1	Uninoculated and unfertilized control	216.8	210.0	202.4	
T2	75% RDF of NPK	204.0	198.0	180.4	
3	100% RDF of NPK	191.4	187.0	173.6	
T4	150% RDF of NPK	156.2	147.4	112.4	
5	100% RDFof NPK through conventional method of applic CD (P>0.05)	ation 176.0 29.11	165.6 28.06	162.0 25.31	

Table 2. Effect of fertigation on soil respiration in maize rhizosphere (COMH 1) under precision farming system

microorganisms. However, increased dosage inhibits the survival of microbes due to osmotic stress created by fertilizers. Among the three doses of fertilizers, plots supplied with 75% RDF of NPK showed maximum *Azospirillum* (0.478 x 10⁵ MPN / g), *Bacillus* (64 x 10⁵ cfu / g) and diazotrophs (132 x 10⁵ cfu / g) population at 28 Days after sowing (DAS), followed by 100% and 150% RDF of NPK (Fig.1).



Fig 1. Effect of fertigation on *Azospirillum*, phosphobacteria and total diazotrophs in the rhizosphere soil of maize (COMH 1) under precision farming system. The treatments ($T_1 - T_5$) are described in Materials and methods.

The results showed that microbial populations were significantly reduced when increasing the dosage of fertilizer. The data from the results suggests that addition of low amount of fertilizer in soil doesn't affect the survival of microorganisms, whereas increased fertilizer dosage leads to accumulation of higher levels of inorganic nutrients in the soil, which inhibits the enzyme activity and survival of microbe's growth. Long term supply of small amount of inorganic amendments in the soil leads to accumulation of more nitrogen nutrients and increases C/N ratio. This leads to slow degradation of organic matter by microorganisms due to decreasing mineralization of organic matter content in soil.

Effect of fertigation on soil respiration

Highest CO₂ evolution (216. 8 mg CO₂ / g / h) was observed in the unfertilized and uninoculated control due to the residual organic matter content of the previous crop, followed by 75% RDF of NPK (204.0 mg CO₂ / g / h) and 100 % RDF of NPK (191.4 mg CO₂ / g / h) during vegetative growth stage (Table 2). The treatment, 150 % RDF of NPK showed

reduction in CO2 evolution (156.6 mg CO2 / g / h) as compared to other treatments due to inhibition of microbial activity, as the result of higher fertilizer dosage. CO2 evolution was maximum during vegetative growth stage (216.8 CO₂ / g / h), which was decreased from flowering (210.0 CO2 / g / h) to harvesting stage (202.4 CO₂ / g / h), due to reduction of active root growth after flowering stage (Table 2). The results support the findings of Wagner and Wolf (1999), who found that organic matter with high C / N ratio is slowly degraded by soil microorganisms. C/N ratio also affects microbial community structure, both prokaryotic (bacterial) and eukaryotic. Sarathchandra et al. (1988) and Perrott et al. (1992) conducted research experiments in temperate regions and the results suggested that, following short term removal of fertilizer application from a grazed and limed upland had no effect on abundance or activity of microorgansims. Lack of effect may be due to part of long history of fertilization and accumulation of nutrient reserves in the soil before the experiment. Therefore, it is likely that effect of removing fertilizers application on soil micro organisms will not be evident until accumulated reserves have been used. So, frequent addition of N fertilizers leads to accumulation of substrate in soil. This inhibits enzyme activities of microbes, which in turn decreases microbial populations also.

Effect of fertigation on soil enzyme activities

There was significant increase in soil enzyme activities *viz.*, urease (78 μ g of NH₄ / g / day), dehydrogenase (110 μ g of TPF /g /day), acid phosphatase (250 μ g of p - nitrophenol / g / h) and alkaline phosphatase (810 μ g of p - nitrophenol / g

/ h) and nitrogenase (1912.30 μ mol of C₂H₄ / g / h) in uninoculated and unfertilized control, followed by 75 % RDF of NPK at 30 DAS (Table 3). Réduction in enzyme activities was observed in 150 % RDF of NPK due to négative impact of fertilizers on survival of microorganisms. This is because of inhibitory

Table 3. Effect of fertigation on soil enzymes activities in maize rhizosphere (COMH 1) under precision farming system

	Treatment	Urease (µg of NH4 /g/day)	Dehydrogenase (µg of TPF /g/ day)	e Nitrogenase (µmoles of ethylene produced / g of soil / h)	Acid Alka phosphatase μ (μg of p - (μg nitrophenol/g/h)	line phosphatase g of p nitrophenol/g/h)
T1	Uninoculated and unfertilized control	78.6	110.0	1912.30	250	810
T ₂	75% RDF of NPK	65.2	86.0	1801.60	180	764
T ₃	100% RDF of NPK	54.4	80.0	1100.00	160	612
T4	150% RDF of NPK	54.0	68.0	952.70	150	584
5	100% RDF of NPK through conventiona method of application	al 55.0	82.6	1653.55	130	684
	CD (P> 0.05)	9.59	13.23	202.844	27.17	107.22

effect of fertilizer N on the nitrogenase activity of diazotrophs. The data showed that the soil enzyme activity was more positively influenced by native soil ecosystem than fertilizer application (Table 3). The current results support the finding of Dick (1988), who has reported that long term application of organic manure increased soil enzyme activity and microbial biomass, but NH4⁺ - N fertilizer caused a decrease of amidase and urease activity related to N fertilizer cycle. Wang (1982) and Zhou (1983) who has reported different type and amount of fertilizers directly affected soil enzymatic activities and then affected nutrient uptake by plant roots.

Conclusion

The results reveal that the maximum population of diazotrophs, *Azospirillum* and phosphobacteria were recorded in 75% RDF through biofertigation of NPK. Maximum soil enzyme activities *viz.*, urease, dehydrogenase, acid and alkaline phosphatase and nitrogenase activities were recorded in the uninoculated and unfertilized control. It is concluded that decreasing the usage of chemical fertilizers and following the organic farming leading to environmental safety and sustainable soil health.

References

- Ahlwaalia, M.S., Singh, B. and Gill, B.S. 1993. Drip irrigation system - its hydraulic performance and influence on tomato and cauliflower crops. *J. Water Manage.*, **1**: 6-9.
- Aref, S. and Wander, M.M. 1998. Long term trends of corn yield and soil organic matter in different crop sequences and soil fertility treatments on the Morrow *Plots. Adv. Agron.*, **62**: 153-161.
- Bauhus, J. and Barthel, R. 1995. Mechanisms for carbon and nutrient release and retention in beech forest gaps. II. The role of soil microbial biomass. *Plant Soil*, 168/169: 585-595.

- Beese, F., Hartmann, A, Beck, T, Rackwitz, R. and Zelles, L. 1994. Microbial community structure and activity in agricultural soils under different management. *Zeitschrift fur pflanzenernahrung* and *Bodenkunde*, **157**: 187-195.
- Bergerson, F.J. 1980. Methods for evaluating Biological nitrogen fixation. John Wiley and Sons., New York. p. 702.
- Brown, G.D. 1973. Mineral nutrition of ectomycorrhizae. In: Marks, G.C., Kozlowsik, T.T. (Eds.), Ectomycorrhiza - Their Ecology and Physiology. Academic Press, New York, 151-205p.
- Dick, R.P. 1988. Influence of long term residue management on soil enzyme activity in relation to soil chemical properties of a wheat fallow system. *Soil Biol. Biochem.*, **6**: 159-164.
- Fauci, M.F. and Dick, R.P. 1994. Soil microbial dynamics: short and long term effects of organic and inorganic nitrogen. *Soil Sci. Soc. Am. J.*, **58**: 801-811.
- Franzluebbers, A.J., Hons, F.M. and Zuberer, D.A. 1995. Soil organic carbon, microbial biomass and mineralisable carbon and nitrogen in sorghum. *Soil Sci. Soc. Am. J.*, **59**: 460 -466.
- Giller, K.E., Beare, M.H., Lavelle, P., Izac, A-M.N. and Swift, M.J. 1997. Agricultural intensification, soil biodiversity and ecosystem function. *Appl. Soil Ecol.*, **6**: 3-16.
- Gunapala, N. and Scow, K. 1998. Dynamics of soil microbial biomass and activity in conventional and organic farming systems. Soil Biol. Biochem., 30: 805-816.
- James, N. 1958. Soil extract in soil microbiology. Can. J. Microbiol., 4: 363-370.
- Jenkinson, D.S. 1988. Determination of microbial biomass carbon and nitrogen levels in soils. In: Wilson, J.R. (Ed.), Soil Biochemistry, CAB International, Wallingford, 368-386p.
- Klein, D.A., Lon, T.C. and Goulding, R.I. 1971. A rapid procedure to evaluate dehydrogenase activity of soils low in organic matter. *Soil Biol. Biochem.*, 3: 385-387.
- Lynch, J.M. and Painting, L.M. 1980. Cultivation and the soil biomass. *Soil Biol. Biochem.*, **12:** 29-33.

- Magdoff, F., Lanyon, L. and Liebhardt, B. 1997. Nutrient cycling, transformations and flows: Implications for a more sustainable agriculture. *Adv. Agron.*, **60**: 1-73.
- Maheswari, R. and Ashok, K.R. and Prahadeeswaran, M. Precision farming Technology Adoption Decisions and Productivity of vegetables in Resource -Poor Environments. Agricultural Economics Research Review, Vol 21 (Conference Number), 2008, 415-424p.
- McLatchey, G.P., Reddy, K.R. 1998. Regulation of organic matter decomposition and nutrient release in a wetland soil. J. Environ. Qual., 27: 1268-1274.
- Okon, Y., Albrect, S.L. and Burris, R.H. 1977. Carbon and ammonia metabolism of Spirillum lipoferum. *J. Bacteriol.*, **128**: 592 -597.
- Parkinson, D., Gray, J.R.G. and Williams, S.T. 1971. In: Methods for studying the Ecology of soil Micro organisms. Oxford Blackwell scientific publications. 116p.
- Perrott, K.W., Sarathchandra, S.U. and Dow, B.W. 1992. Seasonal and fertilizer effect of organic cycle and microbial biomass in the hill country soil under pasture. *Aust. J. Soil Res.*, **30**: 383-394.
- Powlson, D.S., Brookes, P.C and Jenkinson, D.S. 1987. Dynamics of carbon and nitrogen in a long term cropping system and permanent pasture system. *Aust. J. Soil Res.*, **45**: 211-221.
- Pramer, D. and Schmidt, E.L. 1956. Experimental Soil Microbiology. Bulter Publ. Co., Minneapolis, USA, 107p.
- Rennie, 1981. Rennie., R.J. A single medium for the isolation of acetylene reduction (nitrogen fixing) bacteria from soils. *Can. J. Microbiol.*, **27**: 8-14.
- Sarathchandra, S.U., Perrott, K.W, Boase, M.R. and Waller, J.E. 1988. Seasonal changes and the effects of fertilizer on some chemical, biochemical and micro

biological characteristics of high producing pastoral soil. *Biol. Fert. Soils*, **6**: 328-335.

- Saxena, A.K. and Tilak, K.V.B.R. 1998. Free living nitrogen fixers: Its role in crop production. In: Microbes for Health, wealth and sustainable Environment. Ed: Verma, A.K. Malhotra publ. Co, New Delhi. 25 -64.
- Sperber, J.E. 1958. Solubilization of apatite by soil microorganisms producing organic acids. *Aust. J. Agric. Res.*, **9:** 782 -787.
- Tabatabai, M.A. 1982. Soil enzymes, Dehydrogenases. In: Methods of Soil Analysis. Part 2. Chemical and Microbiological properties (Eds R.H. Miller and D.R. Keeney). Agron. Monography, 9, ASA and SSSA, Madison, WI.
- Tabatabai, M.A. and Bremner, J.M. 1972. Assay of urease activity in soils. Soil Biol. Biochem., 4: 479-487.
- Tabatabai, M.A., Fum, M.H. and Basta, N.T. 1992. Effects of cropping systems on nitrification in soils. Comm. *Soil Sci. Plant Anal.*, **23**: 1885 -1891.
- Wagner, G.H. and Wolf, D.C. 1999. Carbon transformations and soil organic matter formation. In: Sylvia, D.M., Fuhrmann, J.J., Hartel, P.G., Zuberer, D.A. (Eds.), Principles and Applications of soil Microbiology, Prentice Hall, New Jersy, 218-258 p.
- Wang, Y.Z. 1982. Significance of several soil enzymatic activities for indicating soil fertility. *Chin. J. Soil Sci.*, **11**: 16-23.
- Wardle, D.A. and Parkinson, D. 1991. A statistical evaluation of equations for predicting total microbial biomass carbon using physiological and biochemical methods. *Agric. Ecosyst. Environ.*, **34**: 75-86.
- Watanabe, I. and Barrauio, W.L. 1980. Low levels of fixed nitrogen for isolation of free living N2 fixing organisms from rice roots. *Nature*, **272**: 256 -266.
- Zhou, L.K. 1983. Role of soil total enzyme activities on evaluating soil fertility. *Acta Pedol. Sinica*,**4**: 413-417.

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