# Effect of Different Organic Crop Rotations on Soil Chemical and Biochemical Properties 

${ }^{1,2}$ Nadia Elabed, ${ }^{1}$ Hanem Grissa, ${ }^{1,2}$ Najla Mousratiand and ${ }^{1}$ Mohamed Ben Kheder<br>${ }^{1}$ Technical Center of Organic Agriculture (CTAB), B.P. 54,<br>${ }^{2}$ High Institute of Agronomy of Chott-Meriem, 4042 Chott Mariem, Sousse, Tunisia


#### Abstract

A long-term study on the effects of different crop rotations on microbial biomass, Dehydrogenase Activity (DHA) and chemical soil properties is reported. The experiment was established in 2000 at the Technical Center of Organic Agriculture Station located in Sousse, Tunisia. The 10 cropping systems ( 8 organic plots and 2 conventional plots as control) were compared. Differences in microbial biomass Carbon (C) and Nitrogen ( N ), dehydrogenase activity, available phosphorus, soil bases, Electrical Conductivity ( EC ) and pH appeared to be related in parts to inputs but perhaps also to differing efficiency of crop rotations at soil fertility maintenance. Overall, the finding indicates that soils in the organic systems had higher microbial biomass, DHA, soil bases, EC and available phosphorus than soils in conventional systems. Organic green house plot, characterized by the most diversified crops ( 19 crops in 10 years), had the highest levels of microbial biomass C and N , dehydrogenase activity, available phosphorus, exchangeable K and Na and $\mathrm{EC} . \mathrm{pH}$ was the highest in an organic open field plot which included a combination of crops (perennial, vegetable and field crops) with different deep rooting.


Key words: Crop rotations, microbial biomass, dehydrogenase activity, chemical soil properties, phosphorus

## INTRODUCTION

Organic farming is a form of agriculture that avoids or largely excludes the use of synthetic fertilizers and pesticides, plant growth regulators and livestock feed additives. An understanding of microbial processes is important for the management of farming systems, particularly those that rely on organic nutrient input (Melero et al., 2006). Microbial processes make a large contribution to the release and availability of nutrients required for crop growth. In organic management systems, Nitrogen ( N ) is supplied in organic form via cover crops and manures and large amounts of Carbon (C) are included in the mass of organic material required to achieve adequate amounts of N (Gunapala and Scow, 1998). Carbon additions of virtually any form to arable soils often increase the amount of microbial biomass (Bohme and Bohme, 2006) and its activity (Shannon et al., 2002; Marinari et al., 2006). Microbial biomass, rather than total organic C has been suggested as a useful and more sensitive measure of change in organic matter status (Melero et al., 2006). Studying microbial biomass C (Cmic) can result in a greater understanding of the biological and chemical changes that occur with different agricultural practices (Anderson and Domsch, 1990). Enzymes may
respond to changes in soil management more quickly than other soil variables and therefore, enzymes might be useful as early indicators of biological change (Bandick and Dick, 1999). Organic manures, such as animal manure, green manure and crop residues, significantly increased the activity of a wide range of soil enzymes, as compared to unamended soil (Martens et al., 1992). A number of studies have shown that organic farming leads to higher soil quality and soil biological activity than conventional farming (Carpenter-Boggs et al., 2000; Fliessbach et al., 2000; Shannon et al., 2002).

The objective of the present study is to evaluate the impact of different organic cropping systems on soil microbial biomass, enzyme activity (dehydrogenase) and chemical properties compared to conventional cropping systems under Mediterranean conditions in Center East of Tunisia.

## MATERIALS AND METHODS

The experiment was carried out at the experimental station of the Technical Center of Organic Agriculture located in Sousse region in the Center East of Tunisia, 35 m above the sea level latitude $35^{\circ} 51^{\prime} 32$ North and longitude $10^{\circ} 35^{\prime} 38^{\prime \prime}$ East Greenwich. The region is

Corresponding Author: Nadia Elabed, Technical Center of Organic Agriculture (CTAB), B.P. 54, 4042 Chott Mariem, Sousse, Tunisia

Agric. J., 9 (1): 51-55, 2014

| Years | Plot 1 | Plot 2 | Plot 3 | Plot 4 | Plot 5 | Plot 6 | Plot 7 | Plot 8 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $2000 \backslash 01$ | Fennel, potato | Potato | Potato | Fallow, potato | Potato | Potato | Pea | Fennel, cabbage |
| $2001 \backslash 02$ | Garlic | Onion | Faba bean | Potato | Potato | Potato | Potato | Artichoke |
| $2002 \backslash 03$ | Potato | Potato | Pea, cauliflower, fennel | Tomato | Tomato, potato | Potato | Tomato, potato | Artichoke |
| $2003 \backslash 04$ | Fallow | Vegetables | Potato | Clover | Clover | Barley | Clover | Artichoke |
| 2004105 | Onion, garlic | Potato | Fallow | Aromatic and medicinal plants | Fennel | Cauliflower, cabbage | Artichoke | Wheat |
| 2005\06 | Potato | Faba bean | Green bean zucchini cucumber (greenhouse) | Aromatic and medicinal plants | Cauliflower, cabbage | Potato | Artichoke | Potato |
| 2006107 | Faba bean | Fennel, leek | Melon, tomato (greenhouse) | Aromatic and medicinal plants | Potato | Pea | Artichoke | Cauliflower |
| $2007 \backslash 08$ | Fennel | Artichoke | Tomato, pepper and eggplant (greenhouse) | Aromatic and medicinal plants | Leek, garlic | Faba bean | Cauliflower, cabbage, potato | Faba bean, maize |
| 2008109 | Bean | Artichoke | Green bean, zucchini (greenhouse) | Aromatic and medicinal plants | Faba bean | Fennel, maize | Faba bean | Pea, potato |
| 2009 110 | Potato | Artichoke | Tomato, green been, zucchini (greenhouse) | Aromatic and medicinal plants | Fennel | Potato | Onion | Cauliflower |

Table 2: Crop rotation of 2 conventional plots

| Years | Plot 9 | Plot 10 |
| :--- | :--- | :--- |
| $2000 \backslash 01$ | Maize (OF) | Artichoke(OF) |
| $2001 \backslash 02$ | Potato (OF) | Fallow (OF) |
| $2002 \backslash 03$ | Fallow (OF) | Potato (OF) |
| $2003 \backslash 04$ | Potato (OF) | Potato (OF) |
| $2004 \backslash 05$ | Pepper (OF) | Oat (OF) |
| $2005 \backslash 06$ | Maize (OF) | Fallow (OF) |
| $2006 \backslash 07$ | Chickpea (OF) | Melon (GH) |
| $2007 \backslash 08$ | Fallow (OF) | Pepper (GH) |
| $2008 \backslash 09$ | Potato (OF) | Tomato (GH) |
| $2009 \backslash 10$ | Potato (OF) | Tomato (GH) |

characterized by a Mediterranean climate with humid mild Winter and hot dry Summer. The yearly precipitations vary from $350-400 \mathrm{~mm}$ and are mainly concentrated between October and April. The annual average temperature ranges from $16-19^{\circ} \mathrm{C}$ with a maximum recorded in July and a minimum one recorded in January.

The soil samples were collected from ten plots (Table 1 and 2). At each sampling plot, 4 representative sub-samples were taken at random from a depth of $5-20 \mathrm{~cm}$. The $0-5 \mathrm{~cm}$ soil layer was discarded to reduce spatial variability and also possible point contamination. The 4 sub-samples were taken with a soil auger, mixed, pooled, giving 40 independent composite samples and transferred in sealed plastic bags to the laboratory. The composite samples were sieved ( $<2 \mathrm{~mm}$ ), homogenized and stored at $4^{\circ} \mathrm{C}$ until the analysis.

Chemical analyses were done according to the Methods of Soil Analysis on air-dried and sieved ( $<2 \mathrm{~mm}$ ) soil samples:

- PH was measured in 1:2.5 soil/water suspension using a glass electrode pH -meter
- Total soluble salts were estimated in the soil saturation extract and were measured by using the electrical conductivity meter (value expressed in $\mathrm{ds} / \mathrm{m}$ ). The soil salinity was evaluated according to the conversion: $1 \mathrm{~m} \mathrm{sec} \mathrm{cm}{ }^{-3}=0.7 \mathrm{~g} \mathrm{~L}^{-1}$
- Exchangeable bases $\left(\mathrm{Na}^{+}, \mathrm{K}^{+} \text {and } \mathrm{Ca}\right)^{++}$were determined after acid digestion with $\mathrm{HF} / \mathrm{HNO}_{3}$ (Liu et al., 2002) by a Beckman single beam flame emission spectrophotometer
- Available phosphorus was determined, according to Olsen method modified and based on the extraction of Phosphor by Sodium Bicarbonate. The dosage was made by spectrophotometer with 660 nm (Pauwels et al., 1992)

Measurement of biomass $\mathbf{C}$ and N : Biomass C was measured by the fumigation incubation method (Jenkinson and Powlson, 1976) from the relationship $\mathrm{Bc}=\mathrm{Fc} / \mathrm{kc}$ where $\mathrm{Fc}=((\mathrm{C}$ in fumigated soil) $-(\mathrm{C}$ in unfumigated soil)) and $\mathrm{kc}=0.45$ (Jenkinson and Ladd, 1981).

Organic C rendered extractable to $0.5 \mathrm{M} \mathrm{K}_{2} \mathrm{SO}_{4}$ by fumigation was measured as described by Jenkinson and Powlson (1976). Briefly, moist soil was exposed to CHCl , for 24 h , the fumigant removed and the soil then extracted with $0.5 \mathrm{M} \mathrm{K}_{2} \mathrm{SO}_{4}$, a non-fumigated control was extracted under the same conditions at the time fumigation started. Organic $C$ in the extracts was determined by dichromate digestion. Total N rendered extractable by fumigation was measured on the same $0.5 \mathrm{M} \mathrm{K}_{2} \mathrm{SO}_{4}$ soil extracts used for measurements of extractable C (Brookes et al., 1985).

The extracts were kept frozen $\left(-18^{\circ} \mathrm{C}\right)$ until analysis. Extracts were analyzed for carbon and nitrogen on a TOC-TNb analyzer (Dimatec, Essen, Germany). Fumigated and unfumigated samples were analyzed close together in order to minimize effects due to time shifts. Standards were measured with the samples as quality controls.

Measurement of Dehydrogenase (DHA): DHA was measured following the procedure described by Alef and Nannipieri (1995). Briefly, DHA was determined using 2,

3,5-Triphenyl-Tetrazoliumchlorid (TTC), as the artificial electron acceptor which is reduced to the red colored 1,3 , 5-Triphenyl-Tetrazoliumformazon (TTF) (Benefield et al., 1977). Then, the formazan was extracted with acetone and measured spectro-photometrically at 546 nm (DHA activity is expressed as $\mu \mathrm{g} \mathrm{TPF} \mathrm{g}{ }^{-1}$ soil $\mathrm{h}^{-1}$ ).

Statistical analysis: Data on all parameters/response variables were subjected to analysis of variance (ANOVA) using SAS. Separation of means was done using multiple range duncan test at $\mathrm{p}=0.05$.

## RESULTS AND DISCUSSION

Figure 1 and 2 show the results of soil pH and Electrical Conductivity (EC). According to these results, the rotation program significantly influenced soil pH and $\mathrm{EC}(\mathrm{p}<0.05)$. Cultivation is likely to increase soil acidity due to increased oxidation of organic matter. The soils experienced are generally alkaline, the pH ranged from 8.08-8.84. The addition of cations via manure application has resulted in higher pH levels in the organic systems. Soil pH in the plot 8 is higher than the other treatments, this plot included perennial plants (artichoke for 3 years), vegetable crops (cabbage, cauliflower, potato and faba bean) and field crops (wheat, maize), this combination of crops with different deep rooting decreased oxidation of organic matter in surface ( $5-20 \mathrm{~cm}$ ) and as result decreased soil acidity. pH is lower than the other treatments in plot 9 (conventional greenhouse), this result can be explained by addition of chemicals fertilizers which are responsible for soil acidity. The pH of the experimental treatments is higher than the optimal value, this may be due to the high pH of the Tunisian soils.

Electrical conductivity is a measure of total cations and anions in solutions and is usually determined largely by Ca and Mg ions. Electrical conductivity levels have been found to be tightly linked to $\mathrm{NO}_{3}$ concentrations in the soil. Nitrification (oxidation of $\mathrm{NH}_{4}-\mathrm{NO}_{3}$ ) acidifies soil, bringing cations into solution. Thus, Ca and Mg concentration in solution and EC levels are highly dependent on N fertility practices. The highest soil EC in this study was obtained in plot 3 which included legumes more than other treatments.

There were significant differences between the levels of soil bases $\left(\mathrm{Ca}^{2+}, \mathrm{K}^{+}, \mathrm{Na}^{+}\right)$measured under different cropping systems (Fig. 3). The general low levels of exchangeable bases is the consequence of crops nutrients uptake in soil and the absence of refund.

Plots 4 and 5 with highest levels of pH had lowest levels of available Phosphorus ( P ). The maximum P availability is when soil pH ranges between 6 and 7 .


Fig. 1: Effect of cropping system on soil pH


Fig. 2: Effect of cropping system on soil Eelectrical Cconductivity (EC)


Fig. 3: Effect of cropping system on soil bases $\left(\mathrm{Na}^{+}, \mathrm{K}^{+}\right.$, $\mathrm{Ca}^{++}$)

Generally, soils with pH of 7.5 and higher have a high calcium concentration that binds P as calcium phosphate creating an insoluble compounds that is not available to plants. Soil available P content in the plot grown after fennel and maize at the last year (plot 5) is lower than the other treatments. Similar results were found by Bahouaoui (2008).

Microbial biomass C varied from 594-150 $\mu \mathrm{g} \mathrm{g}^{-1}$; Microbial biomass N varied from 51-20 $\mu \mathrm{g} \mathrm{g}^{-1}$; dehydrogenase activity varied from $10.95-2.04 \mu \mathrm{~g} \mathrm{TTF} \mathrm{g}{ }^{-1}$ sol $h^{-1}$. Significant differences ( $\mathrm{p}<0.05$ ) were observed between cropping systems. Results of the study showed highest microbial biomass carbon and nitrogen and dehydrogenase activity in the plot number 3. Similar results were obtained by Daniel when tomato crop grown after green bean had a highest microbial biomass carbon and nitrogen in comparison with other cropping systems, this may be due to the high fresh biomass incorporated in the green bean plot before the plantation of main crop. Generally, soil enzyme activities were correlated with microbial biomass, this indicates that enzyme activities were associated with active microorganisms in soil which are the major source of soil


Fig. 4: Effect of cropping system on available phosphorus


Fig. 5: Effect of cropping system on microbiol biomass C


Fig. 6: Effect of cropping system on microbiol biomass N


Fig. 7: Effect of cropping system on dehydrogenase activity
enzymes. In the field trial, the highest values of dehydrogenase activity and microbial biomass C and N contents were most often obtained in the organic soils and the lowest ones occurred in the conventional soils. Enhanced microbial activity in organic farming systems can be attributed mainly to the application of organic manures and higher amounts of diversified crop residues remaining on the fields than in soil under conventional systems (Martyniuk et al., 2001; Mader et al., 1995). The results were the previous of an extra energy source of microbial growth. Moreover, it has been documented that high doses of mineral fertilizers and pesticides, used in
conventional farming systems to protect crops against pests and pathogens, might adversely affect the development and activity of the soil biota (Blevins et al., 1983; Myskow et al., 1996) (Fig. 4-7).

## CONCLUSION

The use of organic farming practices over a 10 years period resulted in higher soil microbial biomass, dehydrogenase activity, available phosphorus and soil bases than the other treatments conducted in conventional farming. This study indicates that the use of animal manure, compost, cover crops, proper crop rotation include a legume and diversified crops and the elimination of synthetic fertilizer results in increased soil fertility.

## ACKNOWLEDGEMENTS

Researchers are thankful to Dr. Jacques Fuchs from Research Institute of Organic Agriculture FIBL Switzerland for microbial biomass and dehydrogenase activity analysis.

## REFERENCES

Alef, K. and P. Nannipieri, 1995. Methods in Applied Soil Microbiology and Biochemistry. Academic Press, London, Pages: 576.
Anderson, T.H. and K.H. Domsch, 1990. Application of eco-physiological quotients ( $\mathrm{qCO}_{2}$ and qD ) on microbial biomasses from soils of different cropping histories. Soil Biol. Biochem., 22: 251-255.
Bahouaoui, M.A., 2008. Effects of different soil building crops and fertilizers on zucchini crop under Mediterranean Organic farming system: Tunisian case. Master Thesis, IAMB, Valenzano, New Jersey.
Bandick, A.K. and R.P. Dick, 1999. Field management effects on soil enzyme activities. Soil Biol. Biochem., 31: 1471-1479.
Benefield, C.B., P.J.A. Howard and D.M. Howard, 1977. The estimation of dehydrogenase activity in soil. Soil Biol. Biochem., 9: 67-70.
Blevins, R.L., M.S. Smith, G.W. Thomas and W.W. Frye, 1983. Influence of conservation tillage on soil properties. J. Soil Water Conservation, 38: 301-305.
Bohme, L. and F. Bohme, 2006. Soil microbiological and biochemical properties affected by plant growth and different long-term fertilization. Eur. J. Soil Biol., 42: 1-12.
Brookes, P.C., A. Landman, G. Pruden and D.S. Jenkinson, 1985. Chloroform fumigation and release of soil nitrogen: A rapid direct extraction method to measure microbial biomass nitrogen in soil. Soil Biol. Biochem., 17: 837-842.

Carpenter-Boggs, L., A.C. Kennedy and S.P. Reganold, 2000. Organic and biodynamic management: Effects on soil biology. Soil Sci. Soc. Am. J., 54: 1651-1659.
Fliessbach, A., P. Mader, D. Dubois and L. Gunst, 2000. Results from a 21 year old field trial. Organic farming enhances soil fertility and biodiversity. FiBL-Dossier No. 1, pp: 15
Gunapala, N. and K.M. Scow, 1998. Dynamics of soil microbial biomass and activity of conventional and organic farming systems. Soil Biol. Biochem., 30: 805-816.
Jenkinson, D.S. and D.S. Powlson, 1976. The effects of biocidal treatments on metabolism in soil-V: A method for measuring soil biomass. Soil Biol. Biochem., 8: 209-213.
Jenkinson, D.S. and J.N. Ladd, 1981. Microbial Biomass in Soil: Measurement and Turnover. In: Soil Biochemistry, Paul, E.A. and J.N. Ladd (Eds.). Marcel Dekker, New York, USA., pp: 415-471.
Liu, F., C. Colombo, P. Adamo, J. Z. He and A. Violante, 2002. Trace elements in manganese-iron nodules from Chinese Alfisol. SSSA J., 66: 661-670.
Mader, P., A. Fiessbach, A. Wiemken and U. Niggli, 1995. Assessement of Soil Microbial Status under Long-Term Low Input (Biological) and High Input (Conventional) Agriculture. In: Effects of Low and High External Input Agriculture on Soil Microbial Biomass and Activities in View of Sustainable Agriculture, Mader, P. and J. Raupp (Eds.). Research Institute of Organic Agriculture, Switzerland, pp: 24-38.

Marinari, S., R. Mancinelli, E. Campiglia and S. Grego, 2006. Chemical and biological indicators of soil quality in organic and conventional farming systems in Central Italy. Ecol. Indicators, 6: 701-711.
Martens, D.A., J.B. Johnson and W.T. Frankenberger, 1992. Production and persistence of soil enzymes with repeated additions of organic residues. Soil Sci., 153: 53-61.
Martyniuk, S., A. Gajda and J. Kus, 2001. Microbiological and biochemical properties of soils under cereals grown in the ecological, conventional and integrated system. Acta Agrophys., 52: 185-192.
Melero, S., J.C.R. Porras, J.F. Herencia and E. Madejon, 2006. Chemical and biochemical properties in a silty loam soil under conventional and organic management. Soil Tillage Res., 90: 162-170.
Myskow, W., A. Stachyra, S. Zieba and D. Maziak, 1996. Microbial activity as an indicator of soil fertility and productivity. Rocz. Glebozn., 47: 89-89.
Pauwels, J., E. van Ranst, M. Verloo and A.M. Ze, 1992. Manuel de laboratoire de pedologie. Mwthodes d'analyses de sols et de plantes, equipement, gestion de stocks de verrerie et de produits chimiques. Publications Agricoles No. 28, AGCD., Bruxelles, Belgium, pp: 180.
Shannon, D., A.M. Sen and D.B. Johnson, 2002. A comparative study of the microbiology of soils managed under organic and conventional regimes. Soil Use Manage., 18: 274-283.

