

Quality and quantity of soil organic matter as affected by the period of organic farming in Sekem farm, Egypt

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Abstract

Sekem commercial organic farm was chosen for the present work; it is located at Belbeis 20 Km northeast of Cairo city which represented sandy soils. Five plots in Sekem farm were chosen to represent different periods of organic farming application, i.e. 0, 12, 15, 18 and 23 yrs. Surface (0-20 cm) and subsurface (20-40 cm) soil samples were collected in both winter and summer season. The collected soil samples were subjected to the dry sieve analysis to determine and separate the dry aggregate size of <0.25, 0.25-0.50, 0.50-1.00 and 1.00-2.00 mm diameter. The distributions of total organic carbon were studied in the whole soil and its aggregate fractions. The data showed that total organic carbon significantly increased by increasing the period of organic farming in the surface samples especially in the longest period of organic farming. Total organic carbon was concentrated in the finest aggregate fraction (<0.25 mm) for both summer and winter seasons and it was also increased by increasing the period of organic farming. The investigation of the humic and fulvic separates using infrared (IR) spectrophotometry, showed the dominance of carboxylic bands in fulvic especially in the subsurface soil samples which indicated its acidic function. Humic separates showed a relative increased in the intensity of aromatic bands as compared to fulvic separates with increasing the period of organic farming.

Keywords: Quality, quantity, organic farming period, soil organic carbon.

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Introduction

Egypt has a well-developed and still rapidly growing organic farming sector. About 26000 hectares are organically cultivated, accounting for 1.0% of total of agricultural area of Egypt. Organic production concentrates on about 500 farming enterprises. Most of these farms are desert farms irrigated from the Nile (IFOAM and FIBL, 2006).

Organic farming relies on building up soil organic carbon through crop rotation, green manure, compost to maintain soil productivity, excluding synthetic agrichemical. Organic farming protect the environment by building soil organic carbon and increase carbon sequestration (Bellows, 2002). Field experiments have shown that soil organic carbon content is controlled by change in management via the annual input of organic manures and wastes and the rate at which it decays (Mikha and Rice, 2004). Soil organic carbon contained a15% liable form while 85% is in the stable humified form (Pettit, 2000). The relative distribution of the soil humified organic forms i.e. humic and fulvic is mainly controlled by the period of organic farming. Consequently, the present in situ work aimed to investigate the effect of the long term organic farming on the distribution of soil organic carbon and their humic and fulvic characteristics using infrared technique in an organic enterprises at Belbies (SEKEM), Egypt.

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Material and Methods

Sekem farm started organic agriculture in 1977. The farm is located in Bilbeis (20 km north east of Cairo, 30°09'09N and 31°25'46E elev. 70 m). This farm was used as a model farm of the sandy soils in Egypt (the clay range was 3.1-10.2 %). The site of Sekem farms were presented in Figure 1. This farm contains 4 regions with different periods of organic farming practices. Organic farming started since 12, 15, 18, and 23 years for S₁, S₂, S₃, and S₄, respectively, with S₀ used as control (where no farming was initiated in this sector of the farm). The 12yr treatment, S₁, is planted with olive trees, for that, samples in this sector were collected from near trees (S_n) and between trees (S_b). Compost was applied to the soil at rate of 20 m³ yr⁻¹ Fd⁻¹.

Sampling was carried out at the end of each growing seasons, in May and October, which represent the end of winter and summer seasons, respectively. Soil samples were collected from both surface soil layer (0–20 cm) and subsurface soil layer (20–40 cm). Each soil sample was a composite of five sub-samples. The soil samples were air-dried at room temperature, grounded to pass through a 2 mm sieve and stored prior to analysis.



Figure 1. Map of Northern Egypt showing the site of Sekem Farm

The collected soil samples were subjected to the determination of dry aggregate distribution, namely < 0.25mm, 0.25-0.50 mm, 0.50-1.00 mm and 1.00-2.00 mm, using mechanical shaker sieves based on the method of Whalen and Chang (2002).

Total organic carbon was determined in the whole soil samples as well as their dry aggregate separates as described by Walkely-Black method (Nelson and Sommers, 1982).

The classical method of fractionating soil humified organic carbon (HOM) is based on the extraction of organic carbon from soil with caustic alkali and the further subdivision of the extracted material by partial precipitation with mineral acids, and by partial solubility in alcohol. The following groups were obtained: the "humic fraction," insoluble in alkali; the "humic acid fraction," soluble in alkali, insoluble in acid; the "fulvic acid fraction," soluble in alkali, soluble in acid (Hayes, 2006).

After the purification, dried humic and fulvic acid were subjected to the FT-IR examinations by a JesCo FTS 460+ single beam using the KBr technique (2 mg sample in 150 mg KBr) (Celi et al., 1997; Eleerbrock et al., 1999) to obtain absorption spectra of both acids extracts (range of wave number 3900-400 cm⁻¹). All spectra were obtained under the same resolution (1cm⁻¹) and measurement conditions.

Results and Discussion

Distribution of soil organic carbon as affected by organic farming periods

Table 1 showed an increase in total organic carbon (TOM) with the increase in cultivation period in Sekem farm for both surface and subsurface layers in winter and summer seasons. The magnitude of TOM increase was much higher in the surface layer, as it reached after 23 yrs of cultivation almost 18 times that in the beginning, while it was only 5 times in the subsurface layer after 23 yrs of cultivation, due to the surface application of organic compost and plant residues. The application of organic manures for ten years caused superior increases in total SOC as compare to the applications of n and P fertilizers (Mikha and Rice, 2004).

Table 1. Total organic carbon changes in surface and subsurface with different periods of organic farming in winter and summer seasons in Sekem farm.

Season	Depth (cm)	S ₀	S ₁	S ₂	S ₃	S ₄
		mg/kg				
Winter	0-20	560	6132	7203	8119	10789
	20-40	734	1162	1630	1348	3956
Summer	0-20	579	5978	7122	5885	10380
	20-40	767	1032	2516	2482	3894

S₀= control; S₁= 12 yrs; S₂= 15 yrs; S₃= 18 yrs and S₄= 23 yrs.

Rochette and Gregorich (1998) stated that at the end of the growing season @ 50% of carbon of the applied organic manure was remained in soil. Soil Organic carbon could be doubled by annual application rate of 2.0 tons carbon from organic amendments (Gerzabek et al., 2001).

Buildup of soil carbon could only be attained when Carbon input is larger than decomposition rate. Soil management processes such as continuous cropping, cover crop and or rotation includes perennial leguminous crops as well as continuous addition of manures certainly guarantee carbon buildup in the soil. Grant et al. (2001) stated that two year rotation was not sufficient to raise SOC in the uppermost topsoil (15 cm depth) in the absence of organic manuring.

Relationship between long term organic farming and TOM was illustrated in Figure 2. Total organic carbon (TOM) was highly significant relationship related with time for Sekem soil (R²=0.996 and 0.908 for winter and summer seasons, respectively). High R² was also observed for the subsurface layer (0.782 and 0.851 under winter and summer seasons, respectively). This relative decrease in R² was due to the lower amount of organic compost and plant residual that could reached the subsurface layer.

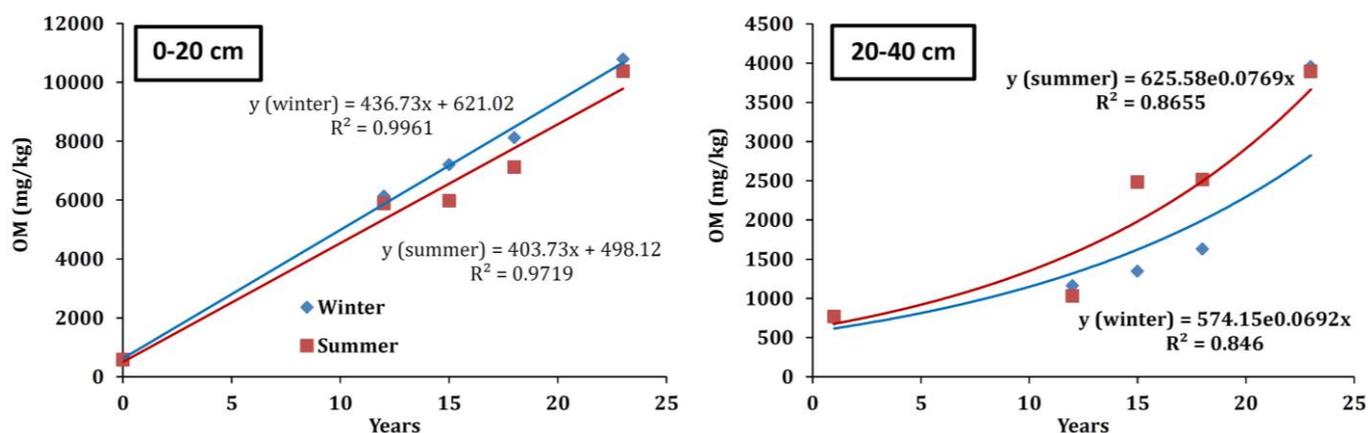


Figure 2. Relationship of OM with time in Sekem farm for winter and summer seasons in surface (0-20 cm) and subsurface (20-40 cm) layers

Distribution of TOM in dry sieve separates with time

Figure 3 represents TOM distribution in dry sieve separates for Sekem farm which indicated that the highest TOM content was recorded in the finest fraction (<0.25 mm) and it increased with increasing period of organic farming. On the other hand, the coarse separates (1.0-2.0 mm) contained the lowest TOM %. This trend was consistently found in winter and summer seasons for the surface and subsurface layers of Sekem farm. Desjardins et al. (1994) confirmed that C_{org} concentrations increased with decreasing soil particle size. In addition, Aoyama et al. (1999) stated that particulate and mineral-associated organic carbon in water-stable aggregates were increased by increasing the rate of organic manure applications.

This could be due to that the smallest size separate has the largest quantity of clay size particles which indicated that organic compounds had been stabilized through the formation of strong clay-organic complexes (Gonzalez and Laird, 2003). In addition, the complexation of light molecular weight (rapid decomposable) organic compounds by clay and silt soil particles also maintain its stability in the soil (Sørensen, 1972). Gerzabek et al. (2001) stated that siltsoil particles had medium capacity to retain soil organic carbon while clay ones slowed down OC turnover rates as it shows the maximum capacity for fixing organic compounds (Buyanovsky et al. 1994).

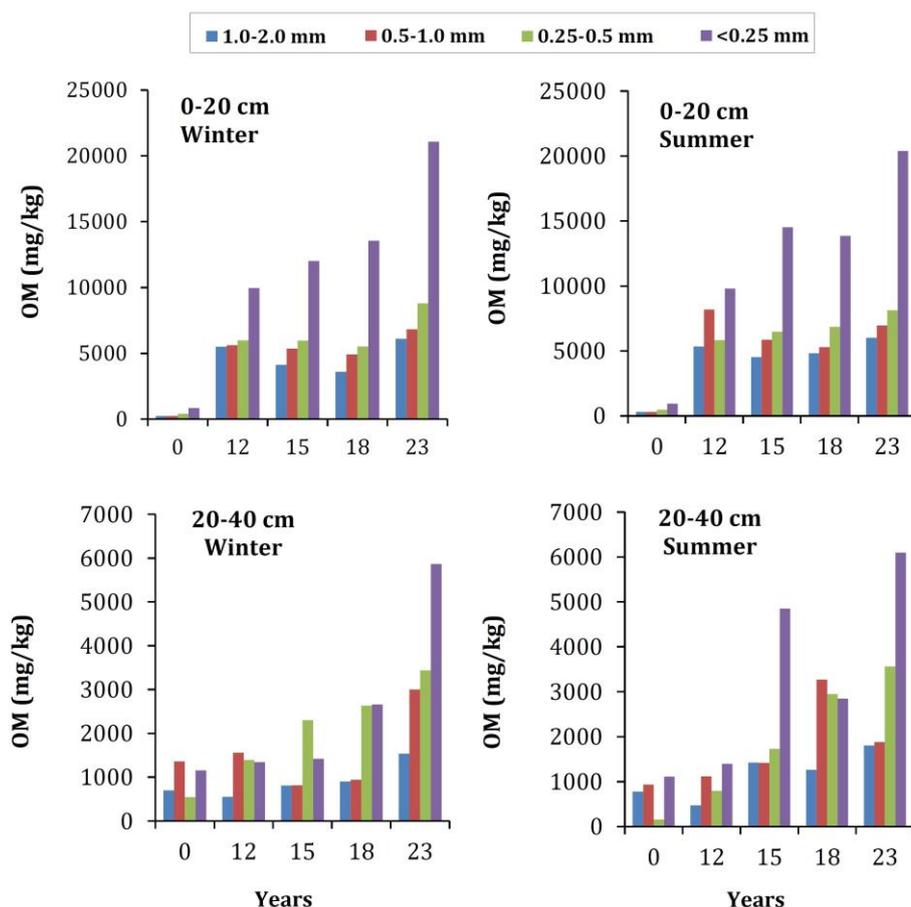


Figure 3. Distribution of OM through dry sieve separates for Sekem in surface and subsurface samples under winter and summer seasons.

Infrared (IR) investigation of humus fractions in Sekem organic farm as affected by organic farming periods

Humic and fulvic acids were separated from the surface (0-20 cm) and subsurface (20-40 cm) soil samples collected from Sekem (23 yrs) and the compost that continuously used as an organic fertilizer. The obtained humic and fulvic acids separates were subjected to IR examination which could be discussed as follows:

IR-characteristics of fulvic acid separate from Sekem farm

Figure 4 represents IR-spectra of fulvic acid separates and the main absorption bands are presented in Table 2. IR-bands were assigned according to [Sánchez-Monedero et al. \(2002\)](#). IR-spectra of fulvic acid exhibit a broad band at 3459-3437 cm^{-1} (stretching of O-H bonds), medium band around 2361-2315 cm^{-1} (stretching of H-N bonds). A well defined band around 1640 cm^{-1} was assigned to aromatic C=C stretching, C=O of ketones and carboxylate ions. A couple of peaks around 1430 cm^{-1} due to aliphatic C-H stretching and peaks around 1030 cm^{-1} due to C-O stretching bond of polysaccharides were also assigned.

Table 2. Corrected relative transmission (T%) for the main IR bands of fulvic acid separates from Sekem organic farm

Bond	compost		Surface (0-20 cm)		Subsurface (20-40 cm)	
	band	T% _{Corr.}	band	T% _{Corr.}	band	T% _{Corr.}
Sekem FA 23 years						
OH st.*	3455	7	3437	2	3439	1
H-N st.	2361	96	2317	90	2318	75
C=C st.	2066	95	2069	85	2087	77
	1868	98	-	-	-	-
C=C	1639	48	1638	25	1638	17
	1517	87	1520	90	-	-
C-H aliph.**	1458	91	1460	85	1459	89
COO st.	1420	92	1400	86	1400	88
KBr	574	55	552	32	572	26

*st.= Stretching band and **aliph. = Aliphatic band.

Relative intensities (transmission T %) of the obtained IR-bands had been corrected to 100% for all IR-spectra in order to make comparisons among them. Table 3 represents the corrected transmission (T %) of IR-bands for compost, surface and subsurface fulvic acid separates. It is well known that the absorbance (A) not transmittance (T %) which expressing the intensity of a specific bond in the examined complex according to Beer-Lambert law (Robincon, 1996). This means that the higher the T% the lower the absorbance and the lower the intensity of the relevant band. Therefore, the obtained intensity differences showed that fulvic acid separates had higher contents of aliphatic and carboxylic H-bonded C=O and amides. The higher content of aliphatic and carboxylic groups in compost fulvic acid separates was widely reported in the literature (Gigliotti et al., 1990), which indicated the higher acidity nature of fulvic acid separates. With respect to the relative transmission (T%) of the IR-bands around 2069 and 1640 cm^{-1} (aliphatic C=C and carboxylic C=O stretching), it was decreased from 96 and 48 % in Sekem compost to 90 and 25 % in the surface layer then reached 75 and 17 %, respectively, in the subsurface layer. This indicated that aliphatic C=C and carboxylic C=O bonds had the lowest absorption intensity in Sekem compost-FA then increased in the surface-FA while the highest intensity of these groups was found in FA separates from subsurface layer. This could be attributed to the progress of FA polymerization which increased double bond intensity through the advancement of the compost decomposition process (Plaza et al., 2007; Abouelwafa et al., 2008). In addition, absorption intensity of carboxylic band was much higher than that of aliphatic C=C bond which indicated the acidic nature of fulvic acid. On the other hand, the intensity around 1520 cm^{-1} (aromatic C=C), 1460 cm^{-1} (aliphatic C-H) and 1420 cm^{-1} (COOH stretching) were slightly changed which indicated that the proportions of these bonds didn't show major differences in Sekem compost and amended soil.

IR-characteristics of humic acid separated from Sekem farm

Figure 5 represents IR-spectra of humic acid separates from Sekem compost, soil surface and subsurface soil layers. IR-spectra of humic acid separates were significantly different from these of FA separates. Hydroxyl O-H stretching was shifted to around 3405 cm^{-1} , new aliphatic COOH group was developed at 2927 cm^{-1} , and new aromatic C-C and C-O-C bonds were developed at 1227 cm^{-1} and 1033 cm^{-1} bands. Additional group of aromatic bands were developed at 1628 cm^{-1} (C=C aromatic), and 1513 and 1415 cm^{-1} (C-H aromatic).

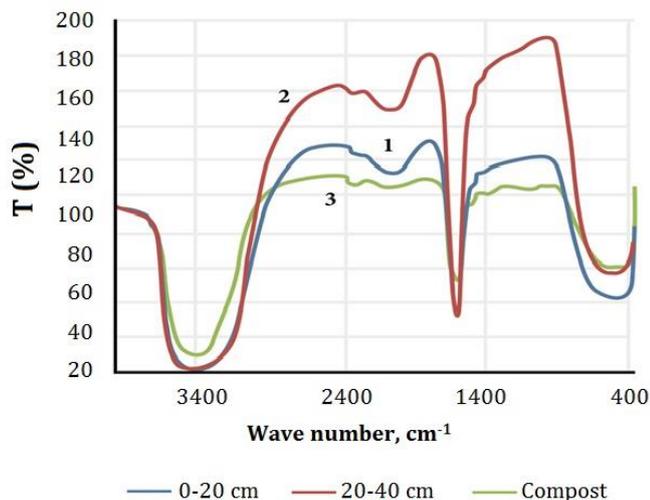


Figure 4. IR-spectra of fulvic acid separates of compost, surface (0-20 cm) and subsurface (20-40 cm) layers of Sekem organic farm cultivated for 23 yrs.

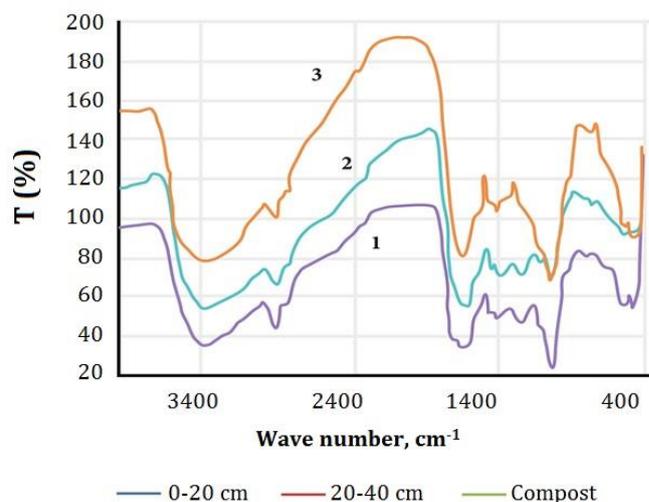


Figure 5. IR-spectra of humic acid separates of compost, surface (0-20 cm) and subsurface (20-40 cm) layers of Sekem organic farm cultivated for 23 yrs.

Regarding the relative intensities of the HA IR-spectra, Table 3 showed a relative increase in the intensity of the aromatic bands (from 16 to 50%) which indicated the abundance of aromatic compounds in humic acid separates as compared to fulvic acid separates which is in agreement with finding of Lguirati et al. (2005) and Guignard et al. (2005).

The relative intensity of OH bond (phenyl and alcohol) was the highest in Sekem compost then decreased in the surface and subsurface soil layer which indicated the polymerization to form humic acid. The relative intensities of HA separates was significantly higher in Sekem compost compared which may confirmed by the high content of stable OM% in Sekem by increasing the period of organic farming.

Table 3. Corrected relative transmission (T%) for the main IR bands of humic acid separates from Sekem organic farm

bond	compost		Surface (0-20 cm)		Subsurface (20-40 cm)	
	band	T% _{Corr.}	band	T% _{Corr.}	band	T% _{Corr.}
Sekem HA 23 years						
OH st.*	3404	14	3406	34	3405	28
Ali. C-H st.	2929	31	2927	42	2927	38
N-H ₂ st.	2367	86	2368	88	-	-
C=N, C=O	-	-	1706	36	-	-
C=C aro.**	1648	16	1628	32	1626	28
	-	-	1513	50	-	-
CH ₂ aliph.	1456	36	1456	48	1455	44
C-H aro.***	1386	34	1415	47	-	-
	-	-	1227	45	1227	42
C-O-C aro.	1031	8	1033	23	1033	42
KBr	522	27	525	52	531	58

* st.= Stretching band; **aliph. = Aliphatic band and *** aro. = Aromatic band

In conclusion, total organic carbon TOM significantly increased by increasing the period of organic farming in the surface samples especially in the longest period of organic farming. Total OM was concentrated in the finest aggregate fraction (<0.25 mm) for both summer and winter seasons and it was also increased by increasing the period of organic farming. The investigation of the humic and fulvic separates using infrared (IR) spectrophotometry, showed the dominance of carboxylic bands in fulvic especially in the subsurface soil samples which indicated its acidic function. Humic separates showed a relative increased in the intensity of aromatic bands as compared to fulvic separates with increasing the period of organic farming.

References

- Abouelwafa, R., Amir, S., Souabi, S., Winterton, P., Ndira, V., Revel, J.C., Hafidi, M., 2008. The fulvic acid fraction as it changes in the mature phase of vegetable oil-mill sludge and domestic waste composting. *Bioresource Technology* 99(14): 6112-6118.
- Aoyama, M., Angers, D.A., N'Dayegamiye, A., 1999. Particulate and mineral-associated organic carbon in water-stable aggregates as affected by mineral fertilizer and manure applications. *Canadian Journal of Soil Science* 79(2): 295-302.
- Bellows, B., 2002. Protecting water quality on organic farms. Available at [Access date: 03.03.2019]: <http://edepot.wur.nl/115572>
- Buyanovsky, G.A., Aslam, M., Wagner, G.H., 1994. Carbon turnover in soil physical fractions. *Soil Science Society of America Journal* 58(4): 1167-1173.
- Celi, L., Schnitzer, M., Nègre, M., 1997. Analysis of carboxyl groups in soil humic acids by a wet chemical method, Fourier-Transform infrared spectrometry and solution-state carbon-13 nuclear magnetic resonance. A comparative study. *Soil Science* 162(3): 189-197.
- Desjardins, T., Andreux, F., Volkoff, B., Cerri, C.C., 1994. Organic carbon and ¹³C contents in soils and soil size-fractions, and their changes due to deforestation and pasture installation in eastern Amazonia. *Geoderma* 61(1-2): 103-118.
- Eleerbrock, R.H., Höhn, A., Rogasik, J., 1999. Functional analysis of soil organic matter as affected by long-term manure treatment. *European Journal of Soil Science* 50(1): 65-71.
- Gerzabek, M.H., Haberhauer, G., Kirchmann, H., 2001. Soil organic matter pools and carbon-13 natural abundances in particle-size fractions of a long-term agricultural field experiment receiving organic amendments. *Soil Science Society of America Journal* 65(2): 352-358.
- Gigliotti, G., Businelli, D., Giusquiani, P.L., 1999. Composition changes of soil humus after massive application of urban waste compost: a comparison between FT-IR spectroscopy and humification parameters. *Nutrient Cycling in Agroecosystems* 55: 23-28.
- Gonzalez, J.M., Laird, D.A., 2003. Carbon sequestration in clay mineral fractions from ¹⁴C-labeled plant residues. *Soil Science Society of America Journal* 67(6): 1715-1720.
- Grant, R.F., Juma, N.G., Robertson, J.A., Izaurralde, R.C., McGill, W.B., 2001. Long-term changes in soil carbon under different fertilizer, manure, and rotation: Testing the mathematical model ecosys. with data from the Breton plots. *Soil Science Society of America Journal* 65(1): 205-214.
- Guignard, C., Lemée, L., Amblès, A., 2005. Lipid constituents of peat humic acids and humin. Distinction from directly extractable bitumen components using TMAH and TEAAc thermochemolysis. *Organic Geochemistry* 36(2): 287-297.

- Hayes, M.H.B., 2006. Solvent systems for the isolation of organic components from soils. *Soil Science Society of America Journal* 70(3): 986–994.
- IFOAM and FiBL, 2006. International Federation of Organic Agriculture Movements (IFOAM), Bonn & Research Institute of Organic Agriculture (FiBL), Frick. The World of Organic Agriculture. Statistics and Emerging Trends. pp. 27–35. Available at [Access date: 03.03.2019]: <https://orgprints.org/5161/>
- Lguirati, A., Ait Baddi, G., El Mousadik, A., Gilard, V., Revel, J.C., Hafidi, M., 2005. Analysis of humic acids from aerated and non-aerated urban landfill composts. *International Biodeterioration & Biodegradation* 56(1): 8-16.
- Mikha, M.M., Rice, C.W., 2004. Tillage and manure effects on soil and aggregate-associated carbon and nitrogen. *Soil Science Society of America Journal* 68(3): 809–816.
- Nelson, D.W., Sommers, L.E., 1982. Total carbon, organic carbon, and organic matter. In: Methods of Soil Analysis, Part 2. Chemical and Microbiological Properties. Page, A.L., Miller, R.H., Keeney, D.R. (Eds.). 2nd Edition. Agronomy Monograph, vol. 9. American Society of Agronomy - Soil Science Society of America, WI, USA. pp. 593–579.
- Pettit, R.E., 2000. Organic carbon, humus, humate, humic acid, fulvic acid, and humin. Available at [Access date: 03.03.2019]: http://www.naturalagsolutions.com/uploads/1/4/7/2/14721182/_organicmatter.pdf
- Plaza, C., Senesi, N., Brunetti, G., Mondelli, D., 2007. Evolution of the fulvic acid fractions during co-composting of olive oil mill wastewater sludge and tree cuttings. *Bioresource Technology* 98(10): 1964-1971.
- Robincon, J.W., 1996. Introduction. In: Atomic Spectroscopy. Robincon, J.W. (Ed.). Second edition, Revised and Expanded. Marcel Dekker, Inc. New York, USA. pp. 1-64.
- Rochette, P., Gregorich, E.G., 1998. Dynamics of soil microbial biomass C, soluble organic C, and CO₂ evolution after three years of manure application. *Canadian Journal of Soil Science* 78(2): 283-290.
- Sánchez-Monedero, M.A., Cegarra, J., García, D., Roig, A., 2002. Chemical and structural evolution of humic acids during organic waste composting. *Biodegradation* 13: 361-371.
- Sørensen, L.H., 1972. Stabilization of newly formed amino-acid metabolites in soil by clay minerals. *Soil Science* 114(1): 5–11.
- Whalen, J.K., Chang, C., 2002. Macroaggregate characteristics in cultivated soils after 25 annual manure applications. *Soil Science Society of America Journal* 66(5): 1637–1647.