

# Monitoring and remediation technologies of organochlorine pesticides in drainage water

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This study was carried out to monitor the presence of organochlorine in drainage water in Kafr-El-Sheikh Governorate, Egypt. Furthermore, to evaluate the efficiencies of different remediation techniques (advanced oxidation processes [AOPs] and bioremediation) for removing the most frequently detected compound (lindane) in drainage water. The results showed the presence of several organochlorine pesticides in all sampling sites. Lindane was detected with high frequency relative to other detected organochlorine in drainage water. Nano photo-Fenton like reagent was the most effective treatment for lindane removal in drainage water. Bioremediation of lindane by effective microorganisms (EMs) removed 100% of the lindane initial concentration. There is no remaining toxicity in lindane contaminated-water after remediation on treated rats relative to control with respect to histopathological changes in liver and kidney. Advanced oxidation processes especially with nanomaterials and bioremediation using effective microorganisms can be regarded as safe and effective remediation technologies of lindane in water.

**Keywords:** lindane, remediation, toxicity, degradation, water.

## INTRODUCTION

One class of organic pollutants which has rightly gained greater attention in environmental studies is the organochlorine pesticides (OCPs). They are highly persistent and toxic in nature, and one of them, dieldrin, has been suspected to be carcinogenic<sup>1</sup>. Due to their persistence in the environment and biological accumulation through the food chain, OCPs can cause environmental damage and affect human health<sup>2-3</sup>. Moreover, due to the limits of water resources in Egypt and the sharp increase of human population, the re-use of drainage water for some purposes (agriculture irrigation and some industrial activities) considered a source of a major concern. However, the re-use of wastewater in agriculture purposes have a great risk on human health in Egypt. Therefore, monitoring of organic pollutants in drainage water and searching for effective remediation technologies to remove these pollutants are in demand to improve the water quality.

Advanced oxidation processes (AOPs), which are constituted by the combination of several oxidants, are characterized by the generation of very reactive and oxidizing free radicals in aqueous solution such as hydroxyl radicals, which possess a great destruction power<sup>4-7</sup>.

Bioremediation considered one of the most environmentally-sound and cost-effective methods for the decontamination and detoxification of pesticides different environmental compartments<sup>8</sup>. The technology of Effective Microorganisms (EMs) was developed during the 1970's at the University of Ryukyus, Okinawa, Japan<sup>9</sup>. Studies have suggested that EMs may have a number of applications, including agriculture, livestock, gardening and landscaping, composting, bioremediation, cleaning septic tanks, algal control and household uses<sup>10</sup>.

However after remediation of pesticide residues in water, toxicity assessment is needed to directly assess the potential hazard of both original pollutants and its metabolites<sup>7</sup>.

In this study the presence of organochlorine pesticides in drainage water in Kafr El-Sheikh governorate was

monitored. The efficiency of advanced oxidation processes with different nano materials and bioremediation with effective microorganisms (EMs) were evaluated to achieve the total degradation of lindane. The histological changes in liver and kidney of rats treated with remediate water relative control were investigated to confirm the complete detoxification of lindane-contaminated water after remediation.

## EXPERIMENTAL

### Chemicals

Organochlorine mixture standard (aldrin, dieldrin, endosulfan, endrin, heptachlor, heptachlor epoxide, lindane, dichlorodiphenyltrichloroethane (DDT), Dichlorodiphenyldichloroethylene (DDE), and dichlorodiphenyldichloroethane (DDD) was obtained from Chem. Service, Inc 660,USA. Lindane with purity of 99.5% was obtained from Central laboratory for Pesticides, Agriculture Research Centre, Giza, Egypt. Zinc oxide (99.99%) and ferric oxide (99.9%) nanoparticles were obtained from Egypt Nanotech Company Limited, El-Wahat road, 6<sup>th</sup> October, Cairo, Egypt. The zinc and ferric oxides particles size are 50 and 40 nm, respectively with a surface area of 60 and 80 m<sup>2</sup>/g, respectively. Hexane and methanol analytical grade solvents were obtained from Sigma – Aldrich Company from Chemicals, U|SA. Hydrogen peroxide and ferric chloride El-Gomhoria Company for Chemical and Glasses, Cairo, Egypt.

### Monitoring of the organochlorine compounds in drainage water

#### Sampling sites

Kafr El-Sheikh (Kotshinar Drainage), Fowa (Fowa Drainage No.11), Metobess (El-Hokss Drainage), Beila (Karakat Drainage), Balteem (Hafeer Shihabeldeen Drainage), Nashart Drainage and El-Hamoul (El-Hamoul Drainage) were selected to be the sampling sites to

cover Kafr El-Sheikh Governorate drainage water areas. These sampling sites were selected according to their proximity to residential areas and agricultural activities. Amber glass bottles were used for sampling and were cleaned by detergent, putted in acid bath, sterilized in the oven and solvent-washed by acetone and hexane before using. Three-liter water samples were collected twice (in the spring and summer) from sampling sites. Glass bottles rinsed twice with the sample water prior to filling and closing. Sampling of water was carried out from the body of running water; the mouth of the bottle was pointed upstream and hands downstream to avoid contamination. The samples were collected for five days during each season. Water samples were acidified with known amount of 1 molar hydrochloric acid to inhibit the biological activity of the possible excite microorganisms. All samples clearly labeled by site number and sampling date. Three replicates were collected from each sampling site and all samples were transferred to the laboratory in ice container for further treatments.

### Extraction procedure

Water samples (500 ml) were extracted twice with 100 ml n-hexane each time. The extracts were combined and filtered through nylon 66 filter (47 mm x 0.45  $\mu\text{m}$ , Supelco, USA). The filtrate was concentrated by rotary evaporation at 50°C to a volume of about 1 mL. For clean up the concentrate was then transferred directly to an activated florisil column, and the OCP fractions were eluted with a mixture of diethyl ether/n-hexane (5:95) at a flow rate of 5 mL/min. Finally, the eluate was concentrated again by rotary evaporation and the final volume of the concentrate was made up to 1.0 mL volume by hexane for the GC-MS analysis<sup>11</sup>. Each sample was extracted and cleaned up three times.

### Gas Chromatography – Mass Spectrometry Analysis

Helium (purity 99.99%) was used as a carrier gas at a constant flow of 1 mL min<sup>-1</sup>. Initial oven temperature was set at 100°C for 2 min, followed by a linear ramp to 180°C at a rate of 5°C min<sup>-1</sup> (hold for 2 min). Subsequently, the temperature was raised to 200°C at a rate of 1.5°C min<sup>-1</sup> followed by a ramp to 250 at a rate of 20°C min<sup>-1</sup>. A final ramp to 280°C was performed at a rate of 30°C and a hold time of 7 min. A split-splitless injector set at 250°C was always used and injections of 1  $\mu\text{L}$  were performed in the splitless mode. Transfer line temperature was set at 285°C and the source temperature at 220°C. The mass spectrometer was operated in the electron impact mode (EI). Electron multiplier voltage was set at 1700 V and the dwell time at 25 ms<sup>12</sup>. The three replicates of each sample were injected to calculate the mean concentration.

### Recovery evaluation

The efficacy of the analytical steps was evaluated by fortifying distilled water samples with the mixture standard of organochlorine pesticides at concentration level of 1 mg L<sup>-1</sup> and then the analytical steps (extraction, cleaning up and determination) that mentioned were performed and replicated three times. Good recovery range (90.8–98.6%) of the tested pesticides was obtained (data not published).

### Photochemical remediation

The scope of the experiments included the following treatments: Nano photo-Fenton-like reagent [Fe<sub>2</sub>O<sub>3</sub>(nano)/H<sub>2</sub>O<sub>2</sub>/UV], nano photo zinc oxide combined with hydrogen peroxide [ZnO (nano)/H<sub>2</sub>O<sub>2</sub>/UV], photo Fenton like reagent (Fe<sup>3+</sup>/H<sub>2</sub>O<sub>2</sub>/UV), and photo zinc oxide combined with hydrogen peroxide (ZnO/H<sub>2</sub>O<sub>2</sub>/UV). For the photo-Fenton-like reagent, a UV mercury lamp model VL-4.LC with a wavelength range of 254 to 365 nm was used for the irradiation of lindane in the aqueous solution. Ferric chloride and ferric oxide nanoparticles were used as sources of the iron catalyst. The solution was prepared by adding a desired amount of lindane (5 mg/L) to filtered El-Hokess drainage (i.e., the highly contaminated site with organochlorine compounds) and carefully agitating the solution. Then, freshly prepared ferric chloride or ferric oxide nanoparticles at a concentration of 50 mg/L as Fe<sup>3+</sup> were added followed by the addition of H<sub>2</sub>O<sub>2</sub> at a concentration of 0.05%. After that, the solution was completed with water up to 1000 mL. The initial pH of the solution was adjusted to 2.8 by using hydrochloric acid 1 M for all experiments<sup>5</sup>. The solution was transferred from the standard flask to a quartz glass cell (1000 mL) and exposed to irradiation of the UV lamp under a constant temperature of 25°C with steering. The solutions (100  $\mu\text{L}$ ) from the irradiated samples were removed at regular intervals (i.e., 10, 20, 40, 80, 160 and 320 min) for high-performance liquid chromatography (HPLC) analysis.

For the ZnO catalyst, 5 mg/L of lindane, with the appropriate amount of ZnO or ZnO nanoparticles (300 mg/L), was shaken carefully before illumination followed by the addition of H<sub>2</sub>O<sub>2</sub> at a concentration of 0.05%. Then, the pH was adjusted to 7, which was the optimum pH for the ZnO catalyst (Derbalah 2009). The suspension was kept in the dark for 30 min before illumination to achieve maximum adsorption of the pesticide onto the semiconductor surface<sup>7</sup>. The solutions from the irradiated samples were removed at regular intervals for HPLC analysis as mentioned before elsewhere. Each experiment was replicated three times for accurate data. Blank experiments were carried out with the tested insecticide alone under the optimum pH and dark conditions were run in parallel at all intervals to assess biotic loss of lindane. The data was negligible due to the high persistence of lindane and the short time.

### HPLC analysis

The irradiated samples were analyzed directly by HPLC (1100 series; Agilent Technologies, Palo Alto, California). The HPLC column used (i.d. of 4.6 mm; length of 250 mm) was filled with Wakosil-II 5 C18-100 (Wako Pure Chemical Industries, Ltd., Osaka, Japan). A mixture of methanol and distilled water (30:70) was used as mobile phase under the isocratic elution mode. The flow rate was maintained at 1 mL/min and the UV detector wavelength was adjusted to be 202 nm<sup>13</sup>.

### Bioremediation technique

The effective microorganisms formulation (EMs1) used for bioremediation of lindane was obtained from the Egyptian Ministry of Agriculture, Giza, Egypt. This formulation contains 60 species of beneficial microorganisms

grown in special media and produced in Egypt under supervision of the Japanese EMRO Scientific Organization (Okinawa, Japan). Enrichment and propagation were carried out in sterilized 250-mL Erlenmeyer flasks using 190 mL mineral salt medium (MSM)[14] and 10 mL of effective microorganisms (5 mL from the formulation) supplemented with lindane at a concentration of 5 mg/L. The cultures were incubated at 30°C, pH 7 and 150 rpm as optimum conditions for the growth of the tested effective microorganisms<sup>15</sup>. Samples were collected at 0, 3, 8, 11, 15, 19, and 23 days for monitoring the degradation of the tested insecticide. Control flasks of equal volumes of mineral salt liquid (MSL) medium and the tested insecticide without the effective microorganisms were run in parallel at all intervals to assess biotic loss. The collected water samples of the tested insecticide were filtered using syringe filter (0.2 mm)<sup>15</sup> followed by HPLC analysis as mentioned before. Each experiment was replicated three times for accurate data.

### Toxicity test

To confirm the complete detoxification of lindane in treated water, toxicity test was conducted on rats. Lindane contaminated-waters after treatment with Fe<sup>3+</sup>/H<sub>2</sub>O<sub>2</sub>/UV, Fe<sub>2</sub>O<sub>3</sub>(nano)/H<sub>2</sub>O<sub>2</sub>/UV, ZnO/H<sub>2</sub>O<sub>2</sub>/UV, ZnO(nano)/H<sub>2</sub>O<sub>2</sub>/UV and EMs were orally administrated to the tested rats. This test was carried out to measure the effect of the possible remaining lindane (parent or metabolites) in the water samples after remediation on rats with respect to histological changes in liver and kidney relative to control.

Adult rats (Sprague dauley) with 100–120 g of weight, obtained from Faculty of Veterinary Medicine, Kafr-El-Sheikh University were used. Rats were housed in polypropylene cages under standard conditions with free

access to drinking water and food. The animals were randomly divided into five groups, each comprising of three animals and water samples (possibly contain lindane or its toxic metabolite) after remediation by different treatments were given to rats as oral administration. Water samples were adjusted to neutral pH, filtered and was free of hydrogen peroxide before orally administrated to rats. Control group rats was fed with normal diet and given oral dose containing no lindane. After 21 days, the rats were scarified under anesthesia and the kidney and liver organs were removed and prepared for histopathological examination according to the method described by Bancroft and Stevens<sup>16</sup>. The histopathology test was carried out at Department of Histopathology, Faculty of Veterinary Medicine, Cairo University Egypt.

## RESULTS AND DISCUSSION

### Monitoring of organochlorine pesticides in drainage water

The analytical parameters of organochlorine pesticides and maximum residue limits of these pesticides are shown in Table (1). The results of wastewater analysis from different sampling sites in Kafr El-sheikh governorate showed the presence of several organochlorine pesticide residues (aldrin, dieldrin, endosulfan, endrin, heptachlor, heptachlor epoxide, lindane, p, p-DDT, p, p-DDE and DDD) in the two sampling times (Tables 2–3). The concentrations of organochlorine pesticides ranged from 0.01 to 0.980 µg L<sup>-1</sup> in drainage water at all sampling sites. With the concerning the sampling sites, the results showed that El-Hokss drainage was the highest contaminated site with organochlorine pesticides while Fowa drainage no. 11 was the lowest contaminated one. With the respect to the detection frequency, lindane was the

**Table 1.** Analytical method parameters of OCPs by the proposed method

Compound name	Limit of detection [ng/L]	Limit of quantification [ng/L]	Correlation coefficients [R <sup>2</sup> ]
Lindane	5.0	20	0.992
Heptachlor	4.0	20	0.995
Aldrin	6.0	40	0.998
Hept. epoxide	15	50	0.994
Endosulfan	14	45	0.998
p,p-DDE	10	75	0.997
Dieldrin	7.0	25	0.983
Endrin	15	100	0.996
DDD	20	60	0.999
p,p-DDT	8.0	20	0.990

**Table 2.** Mean concentration of detected organochlorine pesticides (µg L<sup>-1</sup>) at all sampling sites in spring season

Compounds	Fowa Drainage No.11	El-Hokss Drainage	Karakat Drainage	Hafeer shihab eldeen Drainage	Nashart Drainage	El-Hamoul Drainage	Kotshinar Drainage [Kfs]	Maximum Residue limits [µg/L]
Lindane	0.49 ±0.05	0.98 ±0.1	0.17 ±0.01	0.20 ±0.02	0.95 ±0.08	0.88 ±0.08	0.58 ±0.05	10
Heptachlor	0.19 ±0.01	0.86 ±0.07	0.58 ±0.04	N.D.	0.53 ±0.05	N.D.	0.91 ±0.08	10
Aldrin	0.24 ±0.02	0.96 ±0.09	N.D.	0.88 ±0.07	N.D.	N.D.	N.D.	4.0
Hept. epoxide	N.D.	0.97 ±0.09	0.05 ±0.01	0.15 ±0.01	N.D.	0.12 ±0.01	0.24 ±0.02	10
Endosulfan	N.D.	N.D.	N.D.	0.26 ±0.02	N.D.	0.31 ±0.03	0.18 ±0.01	20
p,p-DDE	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.0
Dieldrin	0.51 ±0.04	0.33 ±0.03	N.D.	0.18 ±0.01	0.14 ±0.01	N.D.	N.D.	4.0
Endrin	N.D.	0.88 ±0.08	0.68 ±0.05	0.65 ±0.05	0.23 ±0.02	0.25 ±0.02	0.27 ±0.02	2.0
DDD	N.D.	0.77 ±0.06	N.D.	N.D.	0.19 ±0.01	N.D.	ND	1.0
p,p-DDT	0.43 ±0.03	0.33 ±0.02	0.06 ±0.01	N.D.	N.D.	0.31 ±0.03	0.15 ±0.01	1.0

N.D. = not detected.  
Hept. = Heptachlore.

**Table 3.** Mean concentration of detected pesticide residues ( $\mu\text{g L}^{-1}$ ) at all sampling sites in summer season

Compounds	Fowa Drainage No.11	El-Hokss Drainage	Karakat Drainage	Hafeer shihab eldeen Drainage	Nashart Drainage	El-Hamoul Drainage	Kotshinar Drainage [Kfs]	Maximum Residue limits [ $\mu\text{g/L}$ ]
Lindane	0.29 $\pm$ 0.02	0.86 $\pm$ 0.08	0.72 $\pm$ 0.07	0.48 $\pm$ 0.04	0.43 $\pm$ 0.04	0.80 $\pm$ 0.08	0.76 $\pm$ 0.07	10
Heptachlor	N.D.	0.03 $\pm$ 0.001	0.02 $\pm$ 0.001	N.D.	N.D.	N.D.	N.D.	10
Aldrin	0.05 $\pm$ 0.001	0.27 $\pm$ 0.02	N.D.	0.14 $\pm$ 0.01	0.05 $\pm$ 0.005	0.05 $\pm$ 0.001	0.07 $\pm$ 0.001	4.0
Hept. epoxide	0.56 $\pm$ 0.05	0.24 $\pm$ 0.02	N.D.	N.D.	0.22 $\pm$ 0.02	0.25 $\pm$ 0.02	0.08 $\pm$ 0.008	10
Endosulfan	N.D.	0.06 $\pm$ 0.001	0.05 $\pm$ 0.001	0.34 $\pm$ 0.03	0.49 $\pm$ 0.04	0.20 $\pm$ 0.02	ND	20
p,p-DDE	N.D.	0.09 $\pm$ 0.001	0.48 $\pm$ 0.04	N.D.	N.D.	N.D.	0.28 $\pm$ 0.02	1.0
Dieldrin	N.D.	0.01 $\pm$ 0.001	0.01 $\pm$ 0.001	N.D.	N.D.	N.D.	N.D.	4.0
Endrin	N.D.	0.05 $\pm$ 0.005	N.D.	0.75 $\pm$ 0.07	0.20 $\pm$ 0.02	0.54 $\pm$ 0.05	0.56 $\pm$ 0.05	2.0
DDD	N.D.	0.41 $\pm$ 0.04	0.08 $\pm$ 0.008	N.D.	0.39 $\pm$ 0.03	0.07 $\pm$ 0.001	N.D.	1.0
p,p-DDT	0.05 $\pm$ 0.001	0.02 $\pm$ 0.002	N.D.	0.01 $\pm$ 0.001	N.D.	N.D.	0.05 $\pm$ 0.001	1.0

highly detected compounds while DDD was the lowest detected one in all sampling sites. The detection frequency and concentration level of the detected pesticides were higher in spring relative to summer season.

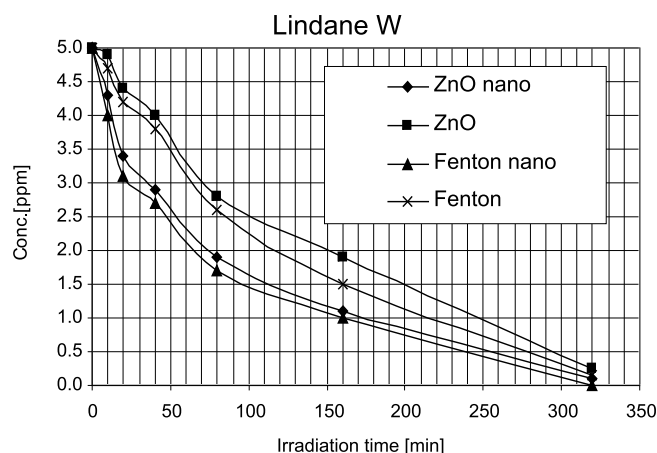
The results of pesticides monitoring showed the presence of several organochlorine compounds in drainage water and this are in agreement with those reported by Abd-Allah and Hesham<sup>17</sup> and Ashry et al.<sup>18</sup>. Spite of some pesticides still present in wastewater after treatment, their concentration level was lower than the maximum residue limits (MRLs) according to Egyptian water quality Standards (Tables 2–3). It is important to note that most of these organochlorines were virtually phased out many years ago and their presence in water residues were from past applications. Firstly, this is attributable to the persistent nature of these compounds. Secondly, water from the Nile originates from the African plateau and crosses eight countries before reaching Egyptian territory (e.g., Sudan, Ethiopia, Uganda, Tanzania, Kenya, Zaire, Rwanda and Burundi). While flowing through these countries, the Nile River is loaded with various types of pesticides and many other contaminants. Thus, it arrives in Egypt after already being contaminated with different pollutants, including the persistent chlorinated pesticides<sup>19</sup>. Thirdly, combustion of domestic wastes is a potential source of PTS in the Egyptian environment with a decreasing abundance in the order PAHs>PCBs>DDTs> HCBs>chlordan>HCHs> endosulfan<sup>20</sup>. Fourthly, Nile River flowing through Kafr El-Zayat City which contained one of the largest pesticides factory in Egypt that flows his drainage contaminated water to Nile water, therefore, the Nile River is loaded with various types of pesticides before reaching Kafr-El-Sheikh Governorate<sup>20</sup>. In addition, organochlorines still have limited use in Egypt as a rodenticide and termiticide<sup>20</sup>.

Finally, the misuse of these pesticides by concerned individuals in addition to the lack of or weak national control is behind the presence of these pesticides in water<sup>18</sup>. The occurrence of such pesticide residues in wastewater represents an environmental and health hazard due to the re-use in agriculture purposes. Frequent monitoring program had urgently needed in order to assess health risks associated with such contaminates especially with chronic exposure or a life-long intake of contaminated drinking water.

#### Degradation of lindane by advanced oxidation processes

The first parameter considered in this study was the losses in lindane concentration with the irradiation time.

The results in Figure 1 showed that, the irradiation under  $\text{Fe}_2\text{O}_3(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  system gave the highest degradation rate of lindane followed by  $\text{ZnO}(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$ ,  $\text{Fe}^{3+}/\text{H}_2\text{O}_2/\text{UV}$  and  $\text{ZnO}/\text{H}_2\text{O}_2/\text{UV}$  systems, respectively. A complete degradation of lindane (100%) was achieved under  $\text{Fe}_2\text{O}_3(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  system followed  $\text{ZnO}(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  (98%),  $\text{Fe}^{3+}/\text{H}_2\text{O}_2/\text{UV}$  (96.8%) and  $\text{ZnO}/\text{H}_2\text{O}_2/\text{UV}$  (95.2%) systems within 320 min of irradiation time, respectively (Fig. 1).

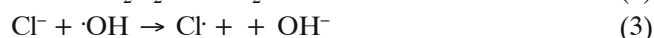


**Figure 1.** Degradation of lindane at initial concentration of 5 mg/L in wastewater  $\text{Fe}_2\text{O}_3(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$ ,  $\text{Fe}^{3+}/\text{H}_2\text{O}_2/\text{UV}$ ,  $\text{ZnO}(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  and  $\text{ZnO}/\text{H}_2\text{O}_2/\text{UV}$  systems

The results showed that, the degradation rate of lindane was enhanced by irradiation under  $\text{Fe}_2\text{O}_3(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  and  $\text{ZnO}(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  systems relative to the degradation under other photochemical remediation systems. This enhancement in lindane degradation rate under  $\text{Fe}_2\text{O}_3(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  and  $\text{ZnO}(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  systems compared to other photochemical irradiation systems may be due to the fact that the stabilized nanoparticles offer much greater surface area and reactivity which lead to higher generation rate of hydroxyl radicals relative to the bulk materials<sup>21–22</sup>.

After 100 min of irradiation time, the degradation rate of the remaining lindane was quite slower than the first 100 min under all photochemical remediation systems. This is might be due to the low remaining concentration of lindane (lower than 20% of its initial concentration) after 100 min of irradiation time which lead to high delivery rate of  $\text{Fe}^{3+}/\text{H}_2\text{O}_2$  and  $\text{ZnO}/\text{H}_2\text{O}_2$  systems corresponds to higher concentrations of these reagents, and this subsequently increase their ability to

compete with lindane as hydroxyl radical scavengers (eqs. 1, 2)<sup>5, 7, 23-25</sup>. Also, chloride and carbonate ions naturally present in water react as hydroxyl radical scavengers<sup>26</sup> as shown in equations 3 and 4.



The degradation rate of lindane under  $\text{Fe}_2\text{O}_3(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  and  $\text{Fe}^{3+}/\text{H}_2\text{O}_2/\text{UV}$  systems was higher than that under  $\text{ZnO}(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  and  $\text{ZnO}/\text{H}_2\text{O}_2/\text{UV}$  systems. This is may be due to the high generation rate of hydroxyl radicals under photo Fenton like reagent (nano or normal) relative to photo zinc oxide combined with hydrogen peroxide (nano or normal)<sup>7</sup>.

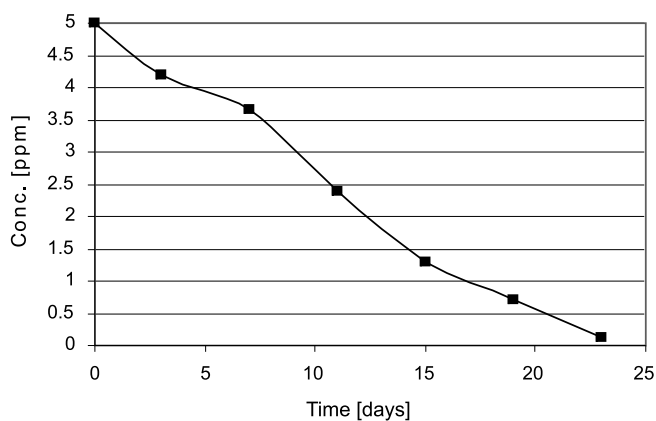
The degradation rate of lindane under  $\text{ZnO}(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  system was higher than that under  $\text{ZnO}/\text{H}_2\text{O}_2/\text{UV}$  system and this is may be due to the effect of particle size of nano zinc oxide. The effect of particle size on the photodegradation efficiency can be ascribed to two reasons. 1) When the size of ZnO crystals decreases, the amount of the dispersion particles per volume in the solution will increase, resulting in the enhancement of the photon absorbance. 2) The surface area of ZnO photocatalyst will increase as the size of ZnO crystals decreases, which will promote the adsorption of more insecticide molecules on the surface<sup>27</sup>.

The degradation rate of lindane under  $\text{Fe}_2\text{O}_3(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  system was higher than that under  $\text{Fe}^{3+}/\text{H}_2\text{O}_2/\text{UV}$  system and this is may be due to the effect of nano ferric oxide particle size which agree with<sup>28-29</sup> who developed a new catalyst using nanosize particles with a high surface area that can accelerate the photo Fenton-like reaction by increasing the hydroxyl radicals generation rate.

The ferric and zinc oxide nanocatalysts are very reactive because the active sites are located on the surface. As such, they have a low diffusional resistance, and are easily accessible to the substrate molecules. Nanocatalysis is but one of the many practical applications of nanotechnology which is concerned with the synthesis and functions of materials at the nanoscale range (lower than 100 nm)<sup>30-32</sup>. An important feature of nanomaterials is that their surface properties can be very different from those shown by their macroscopic or bulk counterparts<sup>33</sup>. As the term suggests, 'nanocatalysis' uses nanoparticles and nanosize porous supports with controlled shapes and sizes<sup>34</sup>. The application of nanoparticles as catalysts of the Fenton-like and photo-Fenton reactions has been described by several investigators<sup>28, 29, 35-37</sup>. In comparison with their microsize counterparts, nanoparticles show higher catalytic activities because of their large specific surface where catalytically active sites are exposed<sup>38</sup>. The advantage of using nanoparticles as catalysts for Fenton-like reagent would more than offset the disadvantage (associated with the use of iron(III) catalysts) of requiring ultraviolet radiation to accelerate the reaction. From all previous discussion, ferric oxide and zinc oxide nanoparticles are potentially useful for remediation of lindane polluted sites<sup>39</sup>.

## Biodegradation of Lindane using effective microorganisms (EMs)

The degradation ability of the effective microorganisms to lindane was illustrated in Figure 2. The effective microorganisms showed high potential in the degradation of the tested insecticide. Nearly 99% of lindane initial concentration (5 mg/L) was degraded within three weeks of incubation with the effective microorganisms.



**Figure 2.** Biodegradation of lindane at initial concentration of 5 mg/L in wastewater by effective microorganisms (EMs)

With the concerning the bioremediation of lindane, effective microorganisms showed high degradation ability against lindane in drainage water. This is may be due to that the effective microorganisms is not one microorganism but a mixture of microorganisms<sup>40</sup> It is also described as a multi-culture of coexisting anaerobic and aerobic beneficial microorganisms<sup>41</sup>. Therefore, its degradation ability to lindane may be faster and effective than using one microorganism. The main species involved in EMs include: lactic acid bacteria (*Lactobacillus plantarum*, *Lactobacillus casei* and *Streptococcus lactis*), photosynthetic bacteria (*Rhodospseudomonas palustris* and *Rhodobacter spaeroides*) yeasts (*Saccharomyces cerevisiae* and *Candida utilis*), actinomycetes (*Streptomyces albus*, and *Streptomyces griseus*) and fermenting fungi (*Aspergillus oryzae* and *Mucor hiemalis*)<sup>42</sup>.

As a conclusion, effective microorganisms could be used in various kinds of aerobic and anaerobic systems for treating agricultural wastes which represent the first point of discharge of many chemicals into environment. The effective and stable degradation capacity of this EMs technology in utilizing and degrading this compound reflected their efficacy in biotechnological application for the bioremediation of such contaminated water. These results indicated that EMs are more stable in retaining their ability to completely degrade lindane because these effective microorganisms live in symbiotic relationships and their influence on the lindane are sum of all activities of these microorganisms. Where the metabolites formed by one type of microorganism may be utilized by other group of microorganisms. This study so far suggested that microorganisms endowed with this property of degradation of toxic pollutants are a boon to mankind. Future studies on the genes responsible for enhanced biodegradation will enable us to elucidate the exact degradation pathway involved in its microbial biodegradation.

## Toxicity assessment

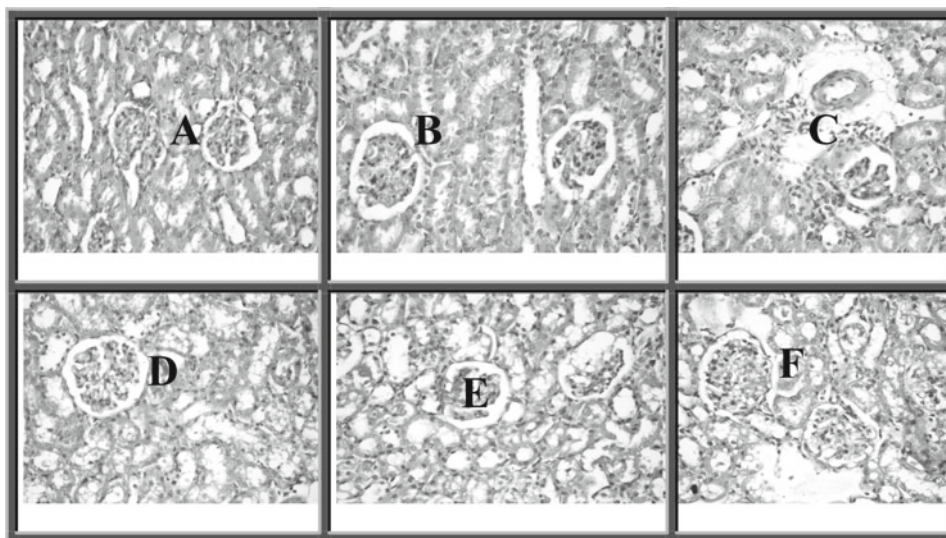
### The histopathological changes in the kidney

The normal structure of kidney tissue is shown in Figure 3A. For the rats treated with lindane after remediation with  $\text{Fe}_2\text{O}_3(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  (Fig. 3B),  $\text{Fe}^{3+}/\text{H}_2\text{O}_2/\text{UV}$  (Fig. 3C),  $\text{ZnO}(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  (Fig. 3D),  $\text{ZnO}/\text{H}_2\text{O}_2/\text{UV}$  (Fig. 3E) and effective microorganisms (Fig. 3F), the tissues were normal like control (Fig. 3B) but for prevascular oedema (Fig. 3C), small vacuulations of epithelial lining renal tubules (Figs. 3D, E) as well as glomerular tufts and epithelial lining renal tubules (Fig. 3F). To confirm the safety of materials used in

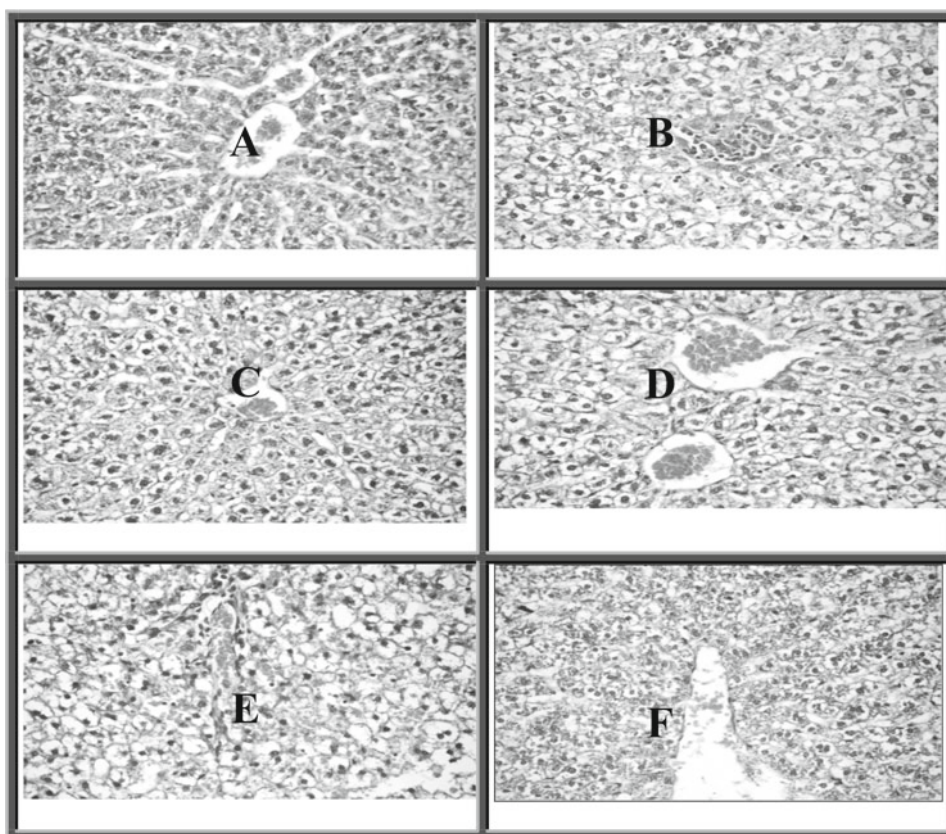
the different remediation processes, some rats treated with  $\text{Fe}_2\text{O}_3(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  (Fig. 3B),  $\text{Fe}^{3+}/\text{H}_2\text{O}_2/\text{UV}$  (Fig. 3C),  $\text{ZnO}(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  (D),  $\text{ZnO}/\text{H}_2\text{O}_2/\text{UV}$  (Fig. 3E) and effective microorganisms (Fig. 3F) without lindane and the kidney tissues were normal like control (data not published).

### The histopathological changes in the liver

The normal structure of liver tissue is shown in Figure 4A. For the rats treated with lindane after remediation with  $\text{Fe}_2\text{O}_3(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  (Fig. 4B),  $\text{Fe}^{3+}/\text{H}_2\text{O}_2/\text{UV}$  (Fig. 4C),  $\text{ZnO}(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  (Fig. 4D),  $\text{ZnO}/\text{H}_2\text{O}_2/\text{UV}$  (Fig. 4E) and Effective microorganisms (Fig. 4F), the tissues were like control but for hydropic



**Figure 3.** Sections in kidney of rats treated with lindane after remediation with  $\text{Fe}_2\text{O}_3(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  (B),  $\text{Fe}^{3+}/\text{H}_2\text{O}_2/\text{UV}$  (C),  $\text{ZnO}(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  (D) and  $\text{ZnO}/\text{H}_2\text{O}_2/\text{UV}$  (E) and Effective microorganisms (F) relative to control (A)



**Figure 4.** Sections in liver of rats treated with lindane after remediation with  $\text{Fe}_2\text{O}_3(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  (B),  $\text{Fe}^{3+}/\text{H}_2\text{O}_2/\text{UV}$  (C),  $\text{ZnO}(\text{nano})/\text{H}_2\text{O}_2/\text{UV}$  (D) and  $\text{ZnO}/\text{H}_2\text{O}_2/\text{UV}$  (E) and Effective microorganisms (F) relative to control (A)

degeneration of hepatocytes (Figs. 4B, C, F), congestion of central vein and hydropic degeneration in hepatocytes (Fig. 4D) and congestion of hepatic sensoids as well as vacuolization of hepatocytes (Fig. 4E). To confirm the safety of materials used in the different remediation processes, some rats treated with Fe<sub>2</sub>O<sub>3</sub>(nano)/H<sub>2</sub>O<sub>2</sub>/UV (Fig. 4B), Fe<sup>3+</sup>/H<sub>2</sub>O<sub>2</sub>/UV (Fig. 4C), ZnO(nano)/H<sub>2</sub>O<sub>2</sub>/UV (Fig. 4D), ZnO/H<sub>2</sub>O<sub>2</sub>/UV (Fig. 4E) and effective microorganisms (F) without lindane and liver tissues were normal like control (data not published).

To evaluate the efficacy of different tested remediation techniques in removing lindane from wastewater, toxicity assessment was carried out with respect to histology test. The histology test for all remediation techniques of lindane in wastewater showed no significant changes in kidney or liver of treated rats relative to control treatment. This implies the complete detoxification of lindane and its possible toxic products in treated wastewater with different remediation techniques. Also, this is implying the safety of all tested chemical and biological remediation techniques on human health especially when we extend the remediation time.

In terms of the removal of used nonmaterials from water after treatment, these could be removed by the addition of natural colloids that make aggregation and sedimentation of these materials and remove them from water<sup>43</sup>. In addition, membranes with suitable porous structures and homogeneous pore-size distribution can separate nanoparticles (NPs) that are less than 10 nm in size. Hence, ultrafiltration and nanofiltration membranes are ideal for separating NPs and large molecules such as proteins because their pore sizes range from 1 to 100 nm<sup>44</sup>. Regarding removal of effective microorganisms, this bioformulation is very safe and has multiple uses such as serving as growth promoters for humans and poultry as well as it used to improve soil fertility etc. Spite of its safety and benefits, it can be remove easy by passing treated water through nanofilter membrane. Since polymer ultra-filtration membranes have been used for the separation of various foods, biological, pharmaceutical systems as well as for water purification<sup>44</sup>.

This study relative to other previous studies used effective microorganisms' formulation for the first time in lindane biodegradation or bioremediation. This is considered a first step for using this safe and effective formulation in the field of wastewater treatment. Moreover, the histology technique to confirm the total detoxification of lindane in remediated water sound interest. Moreover, the toxicity of nanomaterials and effective microorganisms itself was evaluated with respect to histological change in kidney and liver relative to control (data not published) and this also reflect the need to evaluate the side effects or the safety of nanomaterials not only its efficacy.

## CONCLUSIONS

These results clearly indicate the presence of numerous organochlorine compounds in drainage water sampling sites in Kafr El-Sheikh Governorate. The photo-Fenton like reagent and photo zinc oxide combined with hydrogen peroxide showed much promise in the complete

degradation and detoxification of lindane in contaminated drainage water. Effective microorganisms' formulation is promising as effective and safe bioremediation technique for lindane removal in drainage water.

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