

# Evaluation of Microtox and *Daphnia Magna* Acute Toxicity Assays after the Removal of Microorganisms and Organic Compounds from a Pharmaceutical Wastewater using Silver-Loaded Magnetic Nanoparticles

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## Keywords:

Acute toxicity test; *Aliivibrio fischeri* (*Vibrio fischeri*); *Daphnia magna*; Field emission scanning electron microscope (FESEM); Fourier transform infrared spectrophotometer (FTIR); Ibuprofene; Microtox acute toxicity test; Oxytetracycline; pharmaceutical wastewater; silver loaded-magnetic nanoparticles (Ag-Fe<sub>3</sub>O<sub>4</sub> NPs); X-ray diffraction (XRD) analysis; thermogravimetric analysis (TGA).

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## 1. Abstract

**1.1. Background:** In this study, silver-loaded magnetic nanoparticles (Ag-Fe<sub>3</sub>O<sub>4</sub> NPs) were developed under laboratory conditions to treat the secondary settling effluent coming from the biological aerobic activated sludge processes from a Pharmaceutical Wastewater (ww).

**1.2. Material and Methods:** This wastewater contained some pharmaceuticals and bacteria at high concentrations which were not treated at the beginning treatment steps. The effect of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentrations (0.1, 0.5, 1.0, 1.5 mg/l), contacting time (2, 5, 8, 10 and 15 mins), pH levels (4, 7 and 8), power (2, 4 and 6 W/m<sup>2</sup>), frequency (2, 5, 8 kHz). *Salmonella*, *Pseudomonas*, yeast, fungi, total coliforms, fecal coliforms, heterotrophic bacteria and ibuprofene, oxytetracycline removal yields were detected. The Ag-Fe<sub>3</sub>O<sub>4</sub> NPs has a size of 25 nm and the saturation magnetization was recorded as 49 emu/g. The particle shapes and properties were analysed with field emission scanning electron microscope (FESEM) and fourier transform infrared spectrophotometer (FTIR), X-ray diffraction (XRD) and thermogravimetric analysis (TGA). The acute toxicity assays was examined with Microtox (with *Aliivibrio fischeri* also known as *Vibrio fischeri*) and *Daphnia magna* acute toxicity test before and after the removal of microorganisms and organic compounds from a Pharmaceutical ww using Ag-Fe<sub>3</sub>O<sub>4</sub> NPs. ANOVA statistical analysis was used for all

experimental samples. A cost analysis also was performed.

**1.3. Results:** The maximum organism and pollutant yields were around 98% at 0.5 mg/l, after 10 mins at a neutral pH. The recovery of the Ag-Fe<sub>3</sub>O<sub>4</sub> NPs were studied. 89% treatment yields were recorded after 8 months of continuous operation. Adsorptive and photocatalytic studies showed that after a short adsorption time the pollutants and the microorganisms were removed successfully at 2 watt/m<sup>2</sup> power and at 2 kHz frequency. The proposed a novel simple method was applied for the adsorption of ibuprofene and oxytetracycline in water using Ag-Fe<sub>3</sub>O<sub>4</sub> NPs. The TEM image of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs appeared darker than Fe<sub>3</sub>O<sub>4</sub> NPs. Our method of NPs synthesis could not produce uniform particles in size. The XRD patterns of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs showed the 2θ values of 38.11°, 44.32°, 64.24°, 77.61° and 81.57° corresponding to the (111), (200), (220), (311) and (222) planes of cubic Ag, respectively. The Ms values was 61 emu/g for Fe<sub>3</sub>O<sub>4</sub> NPs and 69 emu/g for Ag-Fe<sub>3</sub>O<sub>4</sub> NPs. The increasing Ms values in the nanocomposites (NCs) might have been due to the interactions between the NPs changed the anisotropic energy. 99% maximum ibuprofene adsorption removal yield was obtained at 2.03 mg/g maximum ibuprofene adsorption concentration, at pH = 7.0 and at 21°C, respectively. 99% maximum oxytetracycline adsorption removal yield was observed at 3.04 mg/g maximum oxytetracycline adsorption concentration, at pH = 5.0 and at 21°C, respectively. 93% maximum Ag-Fe<sub>3</sub>O<sub>4</sub>

NPs adsorption yield was measured at 45 min contact time, 7 mg in 500  $\mu$ l of suspension, at 0.1 mg/l Ag-Fe<sub>3</sub>O<sub>4</sub> NPs, at 3.0 mg/l ibuprofene and at 3.0 mg/l oxytetracycline, respectively. 94.44% maximum Microtox acute toxicity yield was observed in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1 g/l after 150 min, at 60°C. However, 90% maximum *Daphnia magna* acute toxicity yield was observed in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1 g/l after 150 min, at 60°C. The removal of microorganisms and organic compounds from a Pharmaceutical ww using Ag-Fe<sub>3</sub>O<sub>4</sub> NPs is a very economical method during different experimental conditions.

**1.4. Conclusions:** The characterization of the adsorbent by means of microscopy, spectroscopy and calorimetry techniques reveal the presence of Ag in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs and the adsorption of ibuprofene and oxytetracycline in Pharmaceutical ww. The adsorption equilibrium is characterized by a Langmuir isotherm model. Microtox acute toxicity test yield is higher than *Daphnia magna* acute toxicity test for the removal of microorganisms and organic compounds from Pharmaceutical ww using Ag-Fe<sub>3</sub>O<sub>4</sub> NPs. Therefore, it can be concluded that the toxicity originating from the Ag-Fe<sub>3</sub>O<sub>4</sub> NPs is not significant and the real acute toxicity throughout the removal of microorganisms and organic compounds was attributed to the Pharmaceutical ww, to the microorganisms and organic compounds in Pharmaceutical ww, to their metabolites and to the disruption by-products. Finally, the removal of microorganisms and organic compounds from a Pharmaceutical ww using Ag-Fe<sub>3</sub>O<sub>4</sub> NPs and the evaluation of Microtox and *Daphnia Magna* acute toxicity assays for these conditions was successfully implemented. The removal of microorganisms and organic compounds from a Pharmaceutical ww using Ag-Fe<sub>3</sub>O<sub>4</sub> NPs is a very economical method during different experimental conditions.

## 2. Introduction

Pharmaceuticals are products used in large doses in daily life considered as contaminants of emerging concern. Due to the large amounts of drugs consumed, the hydrogenic sources suffer from contamination processes that give rise to toxicological effects in humans despite its low concentrations [1, 2]. Many medicines considered as emerging contaminants are constantly detected in groundwater, wastewater treatment plants and water supply. The inefficiency of conventional methods used in water treatment plants to remove the contaminant motivates the development of effective methods to treat effluent contamination [3]. According to the physico-chemical properties of drugs, their degradation products and the characteristics of the soils, these substances can reach the groundwater and contaminate the aquifers or remain retained in the soil, thus affecting the ecosystem and humans through the food chain [4]. Additionally, the portion of medicines not assimilated by the organism, as well as chemical substances administered to animals, usually become part of wastewater. Consequently, different ways of removing medicines in waters have been studied [5]. In recent decades, much attention has been paid to the application

of NPs in environmental purposes. This is because nanosized materials have a large surface area-to-mass ratio and high reactivity [6, 7]. Thanks to these features, NPs have been widely used as catalysts [8], adsorbents [9, 10], detectors [11], and disinfectants [12] in various studies. Due to their non-specific action, NPs as adsorbents are capable of removing a wide variety of contaminants including organics, inorganics, and colloids such as bacterial cells. When making a decrease in microbial contamination is a major goal, one can use some NPs with inherent disinfection capabilities such as magnetic NPs [13]. Therefore, the need for adding other disinfectants such as chlorine (Cl<sup>-1</sup>) and ozone (O<sub>3</sub>), which are associated with adverse side effects, may be eliminated [14].

Magnetite (Fe<sub>3</sub>O<sub>4</sub>) is the strongest magnetic species among the transient metal oxides [15]. The reactive surface of iron-oxides allows for the adsorption of various impurities from water environment [16]. When used as NPs, called Fe<sub>3</sub>O<sub>4</sub> NPs, they release reactive oxygen species (ROS) such as superoxide radicals (O<sub>2</sub><sup>-</sup>), hydroxyl radicals (OH<sup>•</sup>), hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>), and singlet oxygen (<sup>1</sup>O<sub>2</sub>) that can decompose proteins and DNA in bacterial cells [17]. Therefore, Fe<sub>3</sub>O<sub>4</sub> NPs can exert their antibacterial effects through both physical adsorption and chemical disinfection. The feature of Fe<sub>3</sub>O<sub>4</sub> NPs gives them an excellent separation property in the vicinity of an external magnetic field, which is essential for the removal of small-sized NPs from the effluent [14].

The federal government has developed regulations and guidelines to protect people from the possible health effects from long-term exposure to Ag in drinking water [18]. The Environmental Protection Agency (EPA) suggests that the level of Ag in drinking water not be more than 0.05 mg/l. However, in May, 1989, the EPA announced that this restriction on Ag levels in drinking water might be removed. For short-term exposures (1-10 days), EPA suggests that drinking water levels of Ag not be more than 1.142 mg/l. Any release to the environment of more than 453.6 g silver nitrate (AgNO<sub>3</sub>) or 453.6 kg of Ag alone should be reported to the National Response Center. To limit the amount Ag workers are exposed to during an 8-hour shift for a 40-hour work week, the Occupational Safety and Health Administration (OSHA) has set a legal limit (Permissible Exposure Limit or PEL) of 0.01 mg Ag/m<sup>3</sup> air in workroom air. Studies in rats show that drinking water containing very large amounts of Ag (2.6 g Ag/l) is likely to be life-threatening [18].

World Health Organization (WHO) (2002) found that in environments not anthropogenically altered, silver sulfhydrylate (AgHS) or simple sulfur (S) polymer species (HS-Ag-S-Ag-SH) dominate [19]. At higher concentrations, colloidal silver sulfide (Ag<sub>2</sub>S) or silver polysulfide complexes dominate. Under reducing conditions, the Ag ion is sometimes released from the sulfur (S<sub>8</sub>) bearing species. Because concentrations of the Ag ion is typically extremely low in natural environments relative to the available binding sites of S<sub>8</sub>, the Ag ions are quickly combined with other S<sub>8</sub> comp-

lexes. Both scenarios result in essentially non-toxic forms of Ag. Hardness is also a significant control on Ag toxicity. However, it is not the largest control on toxicity as once thought. Below are the data used to create the Environmental Protection Agency (EPA) Ag toxicity equation as a function of hardness, re-evaluated by Hogstrand (1996) [19].

Ag is a noble metal that exhibits a relatively weak oxidation behavior [20]. In large sizes, it is a low-reactive metal. However, in nanoscales, its microbicidal characteristic extremely enhances due to the increased specific surface area [21]. Like  $\text{Fe}_3\text{O}_4$  NPs, Ag NPs can exhibit their antibacterial activity through ROS generation [22]. In addition, other mechanisms have been mentioned such as the interaction of Ag-NPs with surface structures of bacterial cells and the reaction of released Ag ions with  $\text{S}_8$  and phosphorous (P) of cell macromolecules [23]. Accordingly, Ag has been used successfully in previous studies as a bactericidal agent against Gram-negative bacteria, e.g., coliforms [24]. The easy separation of the  $\text{Fe}_3\text{O}_4$  product can facilitate the recovery of invaluable Ag particles. It has been stated that Ag in combination with  $\text{Fe}_3\text{O}_4$  NPs can penetrate biofilms more easily than Ag alone does [21]. Therefore,  $\text{Fe}_3\text{O}_4$ -Ag NPs can show enhanced characteristics for water and wastewater treatment purposes [14].

There are several assumptions for the interaction of Ag and  $\text{Fe}_3\text{O}_4$  NPs. Petrov et al. [25] modified  $\text{Fe}_3\text{O}_4$  NPs with Ag and could detect Ag both around  $\text{Fe}_3\text{O}_4$  NPs and as separate particles in the structure of Ag- $\text{Fe}_3\text{O}_4$  NPs. They also showed that Ag ions replaced  $\text{Fe}^{3+}$  ions in the  $\text{Fe}_3\text{O}_4$  lattice. Therefore, a combination of physical and chemical interactions can play a role in the connection of Ag to  $\text{Fe}_3\text{O}_4$  NPs structure [14].

Various studies in the literature investigated the antibacterial effects of magnetic  $\text{Fe}_3\text{O}_4$  NPs alone or in combination with Ag-NPs as shells. For example, Ghaseminezhad et al. [23] took advantage of the anti-biofilm activity of the compounds against antibiotic-resistant bacterial species in vitro. Joshi et al. [26] employed the composite successfully against *Escherichia coli* (*E. Coli*) using the zone inhibition method on culture plates. However, limited studies are available to compare the effectiveness of  $\text{Fe}_3\text{O}_4$  NPs and Ag- $\text{Fe}_3\text{O}_4$  NPs against bacterial cells in environmental applications. Moreover, the efficacy of the NCs in the decrease of organic content of environmental samples would be interesting to determine. As far as we know, there is no similar study in the literature to investigate the application of  $\text{Fe}_3\text{O}_4$  NPs and Ag- $\text{Fe}_3\text{O}_4$  NPs for advanced purification and disinfection of Wastewater Treatment

Plant (WWTP) effluents.

This study aimed to examine the effectiveness of  $\text{Fe}_3\text{O}_4$  NPs and Ag- $\text{Fe}_3\text{O}_4$  NPs in the removal of Total Coliforms (TC), Fecal Coliforms (FC), Heterotrophic Bacteria (HB), and Chemical Oxygen Demand (COD) from real Pharmaceutical WWTP effluents. In addition to, the particle shapes and properties were analysed with Field Emission Scanning Electron Microscope (FESEM) and fourier transform infrared spectrophotometer (FTIR), X-ray Diffraction (XRD) and Thermogravimetric Analysis (TGA). The acute toxicity assays was examined with Microtox (with *Aliivibrio fischeri* also known as *Vibrio fischeri*) and *Daphnia magna* acute toxicity test before and after the removal of microorganisms and organic compounds from Pharmaceutical ww using Ag- $\text{Fe}_3\text{O}_4$  NPs. ANOVA Statistical Analysis was used for all experimental samples. A cost analysis also was performed.

### 3. Materials and Methods

#### 3.1. Preparation of Ag- $\text{Fe}_3\text{O}_4$ NPs

20 ml of  $\text{H}_2\text{O}$  was heated at  $80^\circ\text{C}$  and continuously stirred under  $\text{N}_2(\text{g})$  atmosphere. Then, 0.56 g  $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$  and 0.2 g  $\text{FeCl}_2 \cdot 4\text{H}_2\text{O}$  were added. When the solids were dissolved, 2 ml of concentrated ammonia ( $\text{NH}_3$ ) solution were incorporated and the solution was stirred for 10 min. The particles were separated using a permanent magnet and the supernatant was discarded. The solid was washed three times with  $\text{H}_2\text{O}$  until the washing liquids were neutral. 0.28 g  $\text{Fe}_3\text{O}_4$  NPs were suspended in 20 ml of  $\text{H}_2\text{O}$ . After, 5.7 ml of diluted 0.011 g/l  $\text{AgNO}_3$  solution were added, the mixture was stirred for 5 min and using the magnet were separated and washed several time with  $\text{H}_2\text{O}$ . Finally, Ag- $\text{Fe}_3\text{O}_4$  NPs were suspended in 20 ml of  $\text{H}_2\text{O}$  again.

#### 3.2. Physicochemical Properties of Ag- $\text{Fe}_3\text{O}_4$ NPs

A Field Emission Scanning Electron Microscopy (FESEM) analysis was performed to check the presence of Ag in the samples. In Figure 1a it is shown the 3D image for  $\text{Fe}_3\text{O}_4$ , while Figure 1b shows the intensity image for Ag- $\text{Fe}_3\text{O}_4$  NPs where the Ag appears as spherical, and shiny due to its high atomic number.

Additionally, energy dispersive X-ray Spectroscopy (EDX) graphs for Ag- $\text{Fe}_3\text{O}_4$  NPs are presented in Figure 2. In the latter, the signal corresponding to Ag appears in the plot. A BET test was carried out to determine the contact surface area of the adsorbent, giving a value equal to  $116.476 \text{ m}^2/\text{g}$ , and a correlation coefficient for the BET isotherm of 0.999.

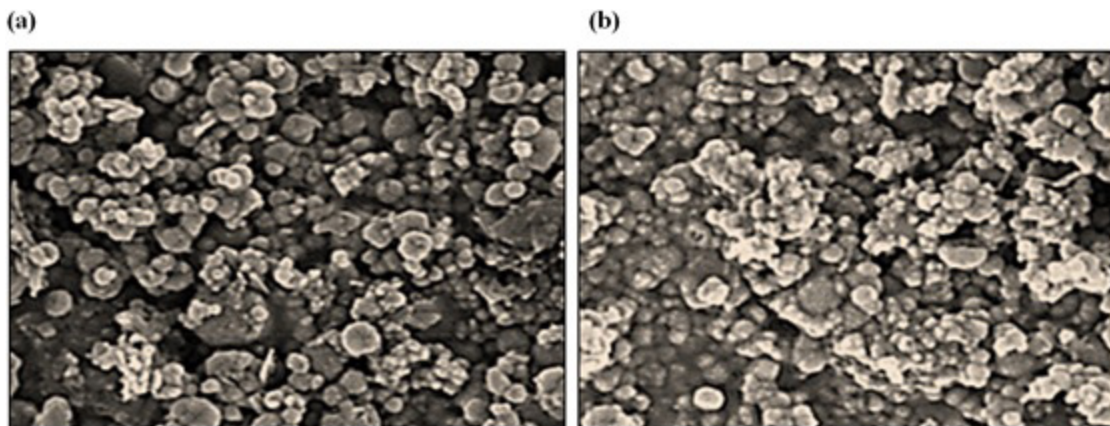


Figure 1: FESEM images of (a)  $\text{Fe}_3\text{O}_4$  NPs and (b)  $\text{Ag-Fe}_3\text{O}_4$  NPs

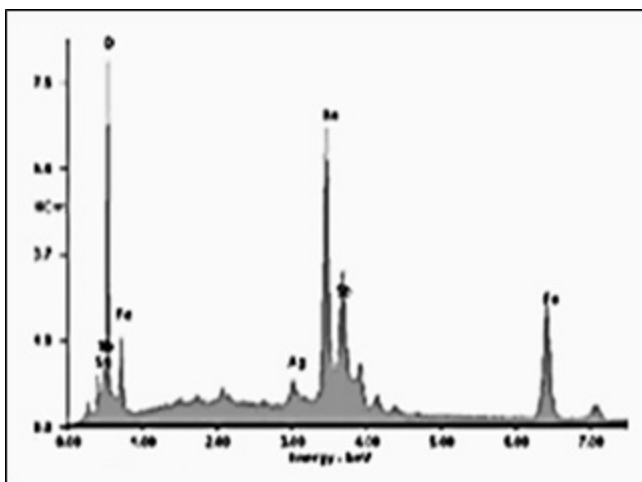


Figure 2: EDX spectra of  $\text{Ag-Fe}_3\text{O}_4$  NPs

### 3.3. Adsorption Studies

A  $\text{H}_2\text{O}$  sample (10 ml) containing pharmaceuticals at concentrations of 0.2 mg/l was placed in a polypropylene tube, 500  $\mu\text{l}$  of  $\text{Ag-Fe}_3\text{O}_4$  NPs suspension were added. After shaking 30 min at  $21^\circ\text{C}$ , the magnet was placed at the bottom of the tube for 5 min and the adsorbent was separated. The supernatant was analyzed for ibuprofene ana oxytetracycline to determine the maximum removal efficiency of the aforementioned chemicals. Figure 3 showed FTIR spectrum for  $\text{Ag-Fe}_3\text{O}_4$  NPs after the adsorption process, where the characteristic IB signals are marked (carbonyl group at  $1704,12 \text{ 1/cm}$  stretch frequencies of Csp<sup>3</sup>-H of isobutyl group at  $2951,93$  and  $2922,15 \text{ 1/cm}$ ; aromatic C=C bond at  $1560,83 \text{ 1/cm}$ ; O-H bond at  $3100 \text{ 1/cm}$ ).

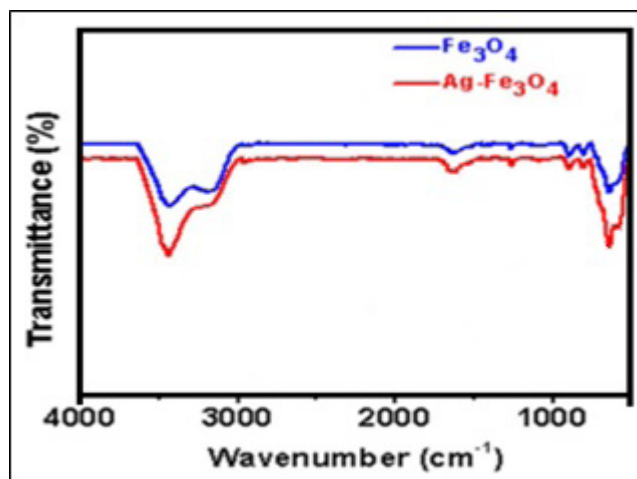


Figure 3: FTIR spectra of  $\text{Fe}_3\text{O}_4$  NPs (blue) and  $\text{Ag-Fe}_3\text{O}_4$  NPs (red)

### 3.4. Acute Toxicity Assays

#### 3.4.1. Microtox Acute Toxicity Test

Toxicity to the bioluminescent organism *Aliivibrio fischeri* (also called *Vibrio fischeri* or *V. fischeri*) was assayed using the Microtox measuring system according to DIN 38412L34, L341, (EPS 1/RM/24 1992). Microtox testing was performed according to the standard procedure recommended by the manufacturer [27]. A specific strain of the marine bacterium, *V. fischeri*-Microtox LCK 491 kit [28] was used for the Microtox acute toxicity assay. Dr. LANGE LUMIX-mini type luminometer was used for the microtox toxicity assay [29].

#### 3.4.2. *Daphnia magna* Acute Toxicity Test

To test toxicity, 24-h born *Daphnia magna* (*D. Magna*) were used as described in Standard Methods section 8711 A, B, C, D, E [30]. After preparing the test solution, experiments were carried out using 5 or 10 *Daphnia magna* introduced into the test vessels. These vessels had 100 ml of effective volume at 7.00–8.00 pH, providing a minimum DO concentration of 6 mg/l at an ambient temperature of 20–25°C. Young *Daphnia magna* were used in the test ( $\leq 24$  h old); 24–48 h exposure is generally accepted as standard for

a *Daphnia magna* acute toxicity test. The results were expressed as mortality percentage of the *Daphnia magna*. Immobile animals were reported as dead *Daphnia magna*.

### 3.5 Statistical Analysis

Multiple regression analysis between y and x variables was performed using the Excell in Windows. The linear correlation was assessed with  $R^2$ . The significance of the correlations between data was determined using the ANOVA Test Statistics [31].

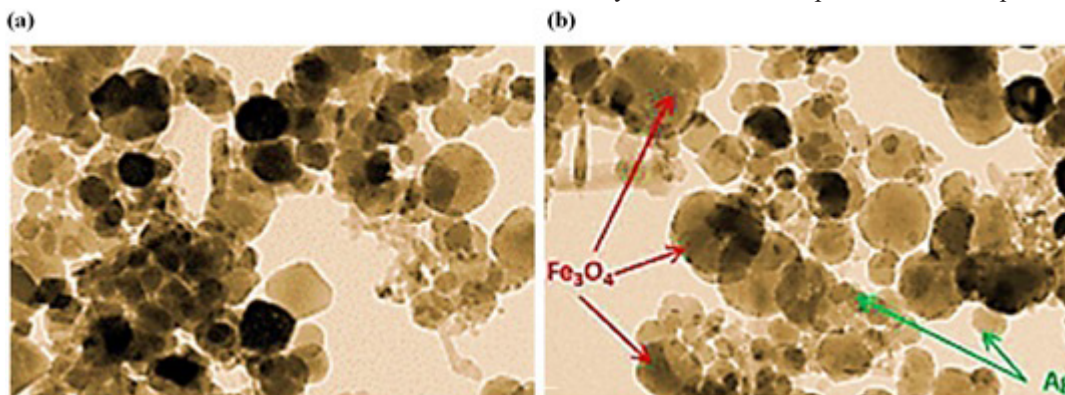
All experiments were carried out three times and the results given as the means of triplicate samplings.

## 4. Results and Discussions

### 4.1. NPs Characteristics

#### 4.1.1. TEM Analysis Results

TEM images were recorded to examine the morphologies of  $\text{Fe}_3\text{O}_4$  NPs and Ag- $\text{Fe}_3\text{O}_4$  NPs and determine the particle sizes. TEM images are presented in Figure 4. As can be observed,  $\text{Fe}_3\text{O}_4$  NPs (Figure 4a) and Ag- $\text{Fe}_3\text{O}_4$  NPs (Figure 4b) were almost semi-spherical in shape. The TEM image of Ag- $\text{Fe}_3\text{O}_4$  NPs appeared darker than that of  $\text{Fe}_3\text{O}_4$  NPs. It is also shown that our method of NPs synthesis could not produce uniform particles in size.



**Figure 4:** TEM images of (a)  $\text{Fe}_3\text{O}_4$  NPs and (b) Ag- $\text{Fe}_3\text{O}_4$  NPs

The size of  $\text{Fe}_3\text{O}_4$  NPs is dependent on various factors including synthesis conditions (e.g. oxygen-free environment) [32], pH of the synthesis medium [33], the ratio of base to iron (Fe) ions, and the length of the alkyl chain, respectively [34]. The mean diameter of particles in our study was 41 and 34 nm for  $\text{Fe}_3\text{O}_4$  NPs and Ag- $\text{Fe}_3\text{O}_4$  NPs, respectively. The reduced size of particles by adding decorating agents has been also observed in the study by Joshi et al. [26]. They used the same method of NPs synthesis as in our study and reported a reduced diameter from 26 to 20 nm after decorating  $\text{Fe}_3\text{O}_4$  NPs with Ag. It seems that a decrease in the amount of iron-oxide precursor per unit volume of the preparation solution might be a reason for the reduced particle size in Ag- $\text{Fe}_3\text{O}_4$  NPs. However, there are reports in the literature [35] showing that the diameter of NPs increases when Ag shells decorated  $\text{Fe}_3\text{O}_4$  cores. The procedure of NPs synthesis has a major contribution to the characteristics of the NCs including the particle size. The particle sizes measured in this study conform to the findings of Naqvi et al. [36] study, reporting the mean size of superparamagnetic iron

oxide NPs as 30 nm, and the findings of Ebrahimi et al. [37] study, producing Ag- $\text{Fe}_3\text{O}_4$  NPs in the range of 23 to 54 nm. The latter study showed that the particle size is dependent on reducing agents used in the synthesis process. Santoyo Salazar et al. [38] conducted a study on  $\text{Fe}_3\text{O}_4$  NPs in the range of 10–40 nm and showed that NPs of smaller than 20 nm had poor  $\text{Fe}_3\text{O}_4$  properties.

#### 4.1.2. XRD Analysis Results

The results of XRD analysis of asprepared NCs (Figure 5). The characterization peaks were observed at  $2\theta$  values of 30.12°, 35.51°, 43.15°, 53.23°, 56.82°, and 62.34°, implying pure  $\text{Fe}_3\text{O}_4$  in the  $\text{Fe}_3\text{O}_4$  NPs structure. These values are in complete agreement with the values reported by Ebrahimi et al. [37] and attribute to the indices (220), (311), (400), (422), (511), and (440) for  $\text{Fe}_3\text{O}_4$  arises, respectively. The XRD patterns of Ag- $\text{Fe}_3\text{O}_4$  NPs showed the  $2\theta$  values of 38.11°, 44.32°, 64.24°, 77.61° and 81.57° corresponding to the (111), (200), (220), (311), and (222) planes of cubic Ag, respectively.

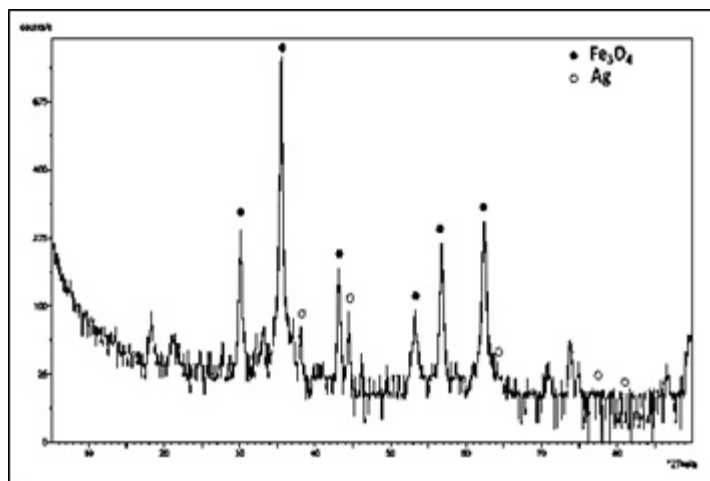


Figure 5: The XRD pattern of  $\text{Fe}_3\text{O}_4$  NPs and Ag NPs.

#### 4.1.3. VSM Analysis Results

The Vibrating Sample Magnetometer (VSM) analysis was conducted to determine the magnetic properties of NPs (Figure 6). Typical superparamagnetic behavior is observed for  $\text{Fe}_3\text{O}_4$  NPs and Ag- $\text{Fe}_3\text{O}_4$  NPs due to the absence of any remanence or coercivity. The analysis showed the saturation magnetization ( $M_s$ ) values of 61 and 69 emu/g for  $\text{Fe}_3\text{O}_4$  NPs and Ag- $\text{Fe}_3\text{O}_4$  NPs, respectively.

The increasing  $M_s$  values in the nanocomposite might have been due to the interactions between the NPs that changed the anisotropic energy. There are conflicting results in the literature regarding the VSM analysis of  $\text{Fe}_3\text{O}_4$ -Ag core-shell structures. Such as, Liu et al. [39], in line with our study, reported an increase in the  $M_s$  values when Ag was doped on the surface of  $\text{Fe}_3\text{O}_4$  NPs, while Li et al. [40] and Ghaseminezhad and Shojaosadati [21].

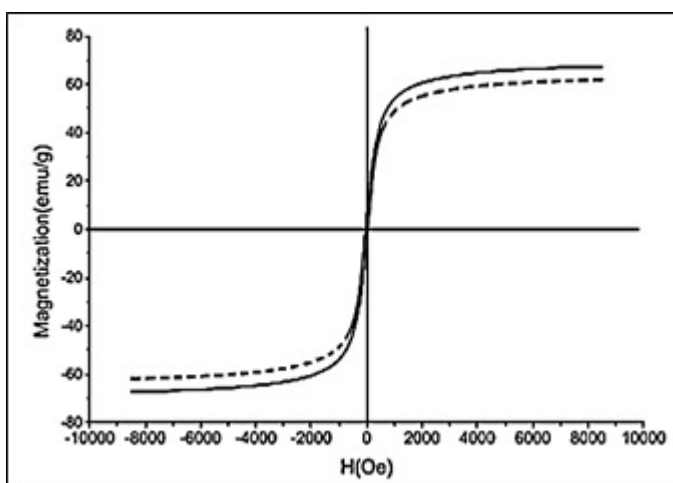


Figure 6: Magnetic hysteresis loops of  $\text{Fe}_3\text{O}_4$  NPs (dashed line) and Ag- $\text{Fe}_3\text{O}_4$  NPs at 21°C.

#### 4.2. Effect of pH on ibuprofene and oxytetracycline adsorption on Ag- $\text{Fe}_3\text{O}_4$ NPs

The effect of pH on the adsorption of ibuprofene and oxytetracycline on Ag- $\text{Fe}_3\text{O}_4$  NPs were studied within the pH ranges varying between 2.0 and 9.0. The highest degree of adsorption was achieved at pH=7.0 (Table 1).

The dependence of adsorption on pH is associated with the point of zero charge (PZC) of the Ag- $\text{Fe}_3\text{O}_4$  NPs and the pKa of the ibuprofene. The PZC is 6.93 for Ag. The ibuprofene is a weak acid (pKa=5.2) and exists as a neutral species por pHpKa41. For pH > 8.0 ibuprofene is deprotonated and Ag surface becomes

negatively charged, thus leading to an electrostatic repulsion which reduces the adsorption efficiency. For pH values greater than pKa but lower than PZC, electrostatic attraction between anionic ibuprofene and the positively charged surface of Ag NPs improves the adsorption capacity. However, the influence of initial pH of oxytetracycline adsorption on Ag- $\text{Fe}_3\text{O}_4$  NPs was different from the ibuprofene. When the pH was lower than 5.0, the adsorption rate increased with pH increasing. When pH was equal to 5.0, the adsorption rate reached the maximum. The maximum removal yields for ibuprofene and oxytetracycline were 99% with maximum adsorption yields of 2.03 mg/g and 3.04 mg/g for ibuprofene and oxytetracycline, respectively.

**Table 1:** Effect of pH on the adsorption yields of 5 mg/l ibuprofene and 5 mg/l oxytetracycline separately in the presence of 3 mg/l Ag-Fe<sub>3</sub>O<sub>4</sub> NPs

pH	Ibuprofene		Oxytetracycline	
	Removal yields (%)	Adsorption (mg/g)	Removal yields (%)	Adsorption (mg/g)
4	67	0.99	56	1.34
5	68	0.99	99	3.04
6	60	0.76	54	1.03
7	99	2.03	50	0.99
8	64	0.97	43	0.76
10	40	0.56	40	0.65

#### 4.3. Effect of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentration on the adsorptions of ibuprofene and oxytetracycline

The Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentrations were increased from 0.1 mg/l up to 6.0 mg/l (Table 2). The ibuprofene and oxytetracycline concentrations were chosen as 3 mg/l at constant concentration. The volume necessary of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs to reach the maximum adsorption efficiency is 500 µl. From 900 µl, the percentage of adsorption decreases due to the high volume of adsorbent in the medium and it is not completely removed with the magnet. 93% maximum Ag-Fe<sub>3</sub>O<sub>4</sub> NPs adsorption yield was measured at 45 min contact time, 7 mg in 500 µl of suspension, at 0.1 mg/l Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentration at 3.0 mg/l ibuprofene and at 3.0 mg/l oxytetracycline, respectively.

#### 4.4. Effect of Contact time on the adsorption of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs to ibuprofene and oxytetracycline

The contact time between Ag-Fe<sub>3</sub>O<sub>4</sub> NPs and the solution containing ibuprofene and oxytetracycline was studied from 1 to 90 min in order to achieve the maximum adsorption efficiency, reached at 45 min. From there, the percentage of adsorption remains constant until 90 min (Table 2).

The proposed method for the adsorption of ibuprofene and oxytetracycline in waters was applied to real water samples partially contaminated with this chemical compound. The concentration of the pollutant in these samples was low or not detected, so they were doped with a known concentrations of ibuprofene (3.0 mg/l) and oxytetracycline (3.0 mg/l) in order to check the viability of the proposed process. Adsorption efficiencies close to 93% were achieved in all cases, thus enabling the method as suitable for its application in real water samples (Table 2).

**Table 2:** Effect of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentration on the adsorption of 3.0 mg/l ibuprofene and 3.0 mg/l oxytetracycline adsorptions

Ag-Fe <sub>3</sub> O <sub>4</sub> NPs concentrations (mg/l)	Ag-Fe <sub>3</sub> O <sub>4</sub> NPs Adsorption (µl)	Contact Time (min)	Ag-Fe <sub>3</sub> O <sub>4</sub> NPs Adsorption efficiencies (%)
0.1	500	45	93
0.5	531	43	92.7
1	580	30	91.2
1.5	612	27	90.9
2	654	24	88.6
2.5	720	10	86.5
3	735	47	83.4
3.5	756	52	71.5
4	783	65	69.8
4.5	800	77	63.4
5	841	81	55.3
5.5	876	88	51.7
6	900	90	48.9

#### 4.5. Desorption and Recycling Studies of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs

The desorption of the ibuprofene and oxytetracycline from the surface of the NPs, at 1 ml HNO<sub>3</sub> solution, at pH=1.0 was used. For that value, interactions between adsorbent and adsorbate are weakened due to the effect of the ionic strength, as stated in the effect of pH on ibuprofene and oxytetracycline adsorption, thus favoring the desorption process. When the ibuprofene and oxytetracycline adsorptions is carried out, after separation from the aqueous solution with the magnet, this solution is decanted. Then, 1 ml HNO<sub>3</sub> solution is added and the mixture is sonicated for 3 min. After, the adsorbent is removed with the magnet and the acid solution is quantified by HPLC, demonstrating that the ibuprofene and oxytetracycline total has been desorbed. Recycling studies

were carried out showing that Ag-Fe<sub>3</sub>O<sub>4</sub> NPs can be used during two additional successive adsorption cycles without losing adsorption capacity. In the fourth adsorption cycle, the capacity decreases down to 89.4%.

#### 4.6. Reuse of the Ag-Fe<sub>3</sub>O<sub>4</sub> Nanocomposites (NCs)

The influence of Ag-Fe<sub>3</sub>O<sub>4</sub> NCs recycling on the degradation rates after one, two and three recycles under UV irradiation were 86%, 86% and 82%, respectively. And the degradation rates of one, two and three recycles under vis light irradiation were 54%, 53% and 50%, respectively. The degradation rate declined slightly with the increasing number of cycles. So Fe<sub>3</sub>O<sub>4</sub>-Ag NCs was effective after three recycling times.

**4.7. Effect of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs Concentrations on the Acute Toxicity Removal Efficiencies in Pharmaceutical ww at Increasing Time and Increasing Temperature**

**4.7.1. Effect of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs Concentrations on the Microtox Acute Toxicity Removal Efficiencies in Pharmaceutical ww at Increasing Time and Increasing Temperature**

The initial EC<sub>90</sub> values at pH=7.0 was found as 600 mg/l at 25°C (Table 3; SET 1). After 60 min, 120 and 150 min, the EC<sub>90</sub> values decreased to EC<sub>60</sub>=612 mg/l to EC<sub>30</sub>=429 mg/l and to EC<sub>10</sub>=350 mg/l in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1 g/l at 30°C (Table 3; SET 3). The toxicity removal efficiencies were 33.33%, 66.67% and 88.89% after 60 min, 120 and 150 min, respectively, in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1 g/l at 30°C (Table 3; SET 3).

The EC<sub>90</sub> values decreased to EC<sub>50</sub>, to EC<sub>20</sub> and to EC<sub>5</sub> after 60 min, 120 and 150 min, respectively, in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs =1 g/l at 60°C (Table 3; SET 3). The EC<sub>50</sub>, the EC<sub>20</sub> and the EC<sub>5</sub> values were measured as 607 mg/l, 482 and 357 mg/l, respectively, in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs =1 g/l at 60°C. The toxicity removal efficiencies were 44.44%, 77.78% and 94.44% after 60 min, 120 and 150 min, respectively, in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1 g/l at 60°C. 94.44% maximum Microtox acute toxicity yield was observed in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1 g/l after 150 min,

at 60°C (Table 3; SET 3).

The EC<sub>90</sub> values decreased to EC<sub>60</sub>=614 mg/l to EC<sub>35</sub>=498 and to EC<sub>15</sub>=354 mg/l after 60 min, 120 and 150 min, respectively, in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs =0.5 g/l at 30°C (Table 3; SET 3). The EC<sub>90</sub> values decreased to EC<sub>65</sub>=603 mg/l to EC<sub>40</sub>=448 and to EC<sub>25</sub>=327 mg/l after 60 min, 120 and 150 min, respectively, in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1.5 g/l at 30°C. The toxicity removal efficiencies were 83.33% and 72.22% in 0.50 and 1.5 g/l Ag-Fe<sub>3</sub>O<sub>4</sub> NPs, respectively, after 150 min at 30°C. It was observed an inhibition effect of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1.5 g/l to *Vibrio Fischeri* microorganisms after 150 min at 30°C (Table 3; SET 3).

The EC<sub>90</sub> values decreased to EC<sub>60</sub>=605 mg/l to EC<sub>30</sub>=494 and to EC<sub>10</sub>=454 mg/l after 60 min, 120 and 150 min, respectively, in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs =0.5 g/l at 60°C (Table 3; SET 3). The EC<sub>90</sub> values decreased to EC<sub>60</sub>=696 mg/l to EC<sub>35</sub>=579 and to EC<sub>20</sub>=452 mg/l after 60 min, 120 and 150 min, respectively, in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs =1.5 g/l at 60°C. The toxicity removal efficiencies were 88.89% and 77.78% in 0.5 and 1.5 g/l Ag-Fe<sub>3</sub>O<sub>4</sub> NPs, respectively, after 150 min, at 60°C. It was observed an inhibition effect of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1.5 g/l to *Vibrio Fischeri* microorganisms after 150 min at 60°C (Table 3; SET 3).

**Table 3:** Effect of increasing Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentrations on Microtox acute toxicity in Pharmaceutical ww at 30°C and at 60°C.

No	Parameters	Microtox Acute Toxicity Values, * EC (mg/l)							
		25°C							
		0. min		60. min		120. min		150. min	
*EC <sub>90</sub>		*EC		*EC		*EC			
1	Raw ww, control	600		EC <sub>80</sub> =610		EC <sub>60</sub> =510		EC <sub>50</sub> =525	
		30°C				60°C			
		0 min	60. min	120 min	150. min	0 min	60. min	120. min	150. min
		*EC <sub>90</sub>	*EC	*EC	*EC	*EC <sub>90</sub>	*EC	*EC	*EC
2	Raw ww, control	600	EC <sub>70</sub> =610	EC <sub>50</sub> =590	EC <sub>40</sub> =680	600	EC <sub>60</sub> =680	EC <sub>40</sub> =610	EC <sub>30</sub> =410
3	Ag-Fe <sub>3</sub> O <sub>4</sub> NPs=0.5 g/l	600	EC <sub>60</sub> =614	EC <sub>35</sub> =498	EC <sub>15</sub> =354	600	EC <sub>60</sub> =605	EC <sub>30</sub> =494	EC <sub>10</sub> =454
	Ag-Fe <sub>3</sub> O <sub>4</sub> NPs=1 g/l	600	EC <sub>60</sub> =612	EC <sub>30</sub> =429	EC <sub>10</sub> =350	600	EC <sub>50</sub> =607	EC <sub>20</sub> =482	EC <sub>5</sub> =357
	Ag-Fe <sub>3</sub> O <sub>4</sub> NPs=1.5 g/l	600	EC <sub>65</sub> =603	EC <sub>40</sub> =448	EC <sub>25</sub> =327	600	EC <sub>60</sub> =696	EC <sub>35</sub> =579	EC <sub>20</sub> =452

\* EC values were calculated based on COD<sub>dis</sub> (mg/l).

**4.7.2. Effect of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs Concentrations on the *Daphnia magna* Acute Toxicity Removal Efficiencies in Pharmaceutical ww at Increasing Time and Increasing Temperature**

As seen in Table 4; SET 1, the initial EC<sub>50</sub> values were observed as 618 mg/l at 25°C. After 60 min, 120 and 150 min, the EC<sub>50</sub> values decreased to EC<sub>35</sub>=325 mg/l to EC<sub>20</sub>=170 mg/l and to EC<sub>10</sub>=70 mg/l in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1 g/l at 30°C (Table 4; SET 3). The toxicity removal efficiencies were 30%, 60% and 80% after 60 min, 120 and 150 min, respectively, in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1.00 g/l at 30°C (Table 4; SET 3).

The EC<sub>50</sub> values decreased to EC<sub>30</sub>=360 mg/l to EC<sub>15</sub>=140 mg/l and to EC<sub>5</sub>=39 mg/l after 60 min, 120 and 150 min, respectively, in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1 g/l at 60°C (Table 4; SET 3). The toxicity removal efficiencies were 40%, 70% and 90% after 60 min, 120 and 150

min, respectively, in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1 g/l at 60°C. 90% maximum *Daphnia magna* acute toxicity yield was observed in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1 g/l after 150 min, at 60°C (Table 4; SET 3).

The EC<sub>50</sub> values decreased to EC<sub>40</sub>=485 mg/l to EC<sub>25</sub>=370 and to EC<sub>15</sub>=190 mg/l after 60 min, 120 and 150 min, respectively, in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=0.5 g/l at 30°C (Table 4; SET 3). The EC<sub>50</sub> values decreased to EC<sub>45</sub>=330 mg/l to EC<sub>30</sub>=170 and to EC<sub>20</sub>=6.50 mg/l after 60 min, 120 and 150 min, respectively, in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1.5 g/l at 30°C. The *Daphnia magna* acute toxicity yields were 70% and 60% in 0.5 and 1.5 g/l Ag-Fe<sub>3</sub>O<sub>4</sub> NPs, respectively, after 150 min, at 30°C. It was observed an inhibition effect of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1.5 g/l to *Daphnia magna* after 150 min, at 30°C (Table 4; SET 3).

The EC<sub>50</sub> values decreased to EC<sub>35</sub>=375 mg/l to EC<sub>20</sub>=23 and to EC<sub>10</sub>=135 mg/l after 60 min, 120 and 150 min, respectively, in



Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=0.5 g/l at 60°C (Table 4; SET 3). The EC<sub>50</sub> values decreased to EC<sub>40</sub>=400 mg/l to EC<sub>25</sub>=125 and to EC<sub>15</sub>=33 mg/l after 60 min, 120 and 150 min, respectively, in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1.5 g/l at 60°C. The Microtox acute toxicity yields were 80% and 70% in 0.5 and 1.5 g/l Ag-Fe<sub>3</sub>O<sub>4</sub> NPs, respectively, after 150 min, at 60°C. It was obtained an inhibition effect of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1.5 g/l to *Daphnia magna* after 150 min, at 60°C (Table 4; SET 3).

The maximum *Daphnia magna* acute toxicity removal was 90% at the Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentration of 1 g/l at 60°C after 150 min (Table 4; SET 3). In this acute toxicity reduction the EC<sub>50</sub> value of raw Pharmaceutical ww decreased to EC<sub>5</sub>=39 mg/l. Low acute toxicity removals found at high Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentrations could be attributed to their detrimental effect on the *Daphnia magna* cells. High Ag-Fe<sub>3</sub>O<sub>4</sub> NPs (> 1.5 g/l) caused lysis in *Daphnia magna* cells. The high salt concentrations caused turgor in *Daphnia magna* cells by increasing the osmotic pressure in the test medium.

A strong significant correlation between EC<sub>50</sub> values and the removal of microorganisms and organic compounds from Pharmaceutical ww showed that the *Daphnia magna* acute toxicity test alone can be considered a reliable indicator of Pharmaceutical ww toxicity (R<sup>2</sup>=0.86, F=4.78, p=0.001). Similarly, a strong linear correla-

tion between threshold concentrations of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs decrease in inhibitions was observed (R<sup>2</sup>=0.87, F=16.72, p=0.001) while the correlation between the inhibition decrease and Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentrations above the threshold values was weak and not significant (R<sup>2</sup>=0.31, F=3.42, p=0.001).

The toxicity of Pharmaceutical ww samples to the tested species prior and after Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentrations. The raw Pharmaceutical ww samples induced 95% motility inhibition to *D. magna* (Table 4; SET 3). This inhibition could be attributed to the mixed recalcitrant carcinogenic Pharmaceutical ww with high benzene rings and to the synergistic effects of the aforementioned more hydrophobic organic compounds with less hydrophobic organic compounds in Pharmaceutical ww. When *D. magnas* were exposed to the effluent Pharmaceutical ww samples after Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentrations of there was a significant reduction in inhibition (from 98.90% to 99.99%) for the acute toxicity after 150 min, at 60°C (Table 4; SET 3).

Low acute toxicity removals found at high Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentrations could be attributed to their detrimental effect on the *D. magna* cells. High Ag-Fe<sub>3</sub>O<sub>4</sub> NPs caused lysis in *D. magna* cells. The high salt concentrations caused plasmolysis in *D. magna* cells by increasing the osmotic pressure in the test medium.

**Table 4:** Effect of increasing Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentrations on *Daphnia magna* acute toxicity in Pharmaceutical ww at 30°C and at 60°C.

No	Parameters	<i>Daphnia magna</i> Acute Toxicity Values, * EC (mg/l)							
		25°C							
		0. min		60. min		120. min		150. min	
*EC <sub>50</sub>		*EC		*EC		*EC			
1	Raw ww, control	618		EC <sub>45</sub> =425		EC <sub>40</sub> =325		EC <sub>35</sub> =182	
		30°C				60°C			
		0 min	60. min	120. min	150. min	0. min	60 min	120. min	150. min
		*EC <sub>50</sub>	*EC	*EC	*EC	*EC <sub>50</sub>	*EC	*EC	*EC
2	Raw ww, control	618	EC <sub>40</sub> = 510	EC <sub>35</sub> = 340	EC <sub>30</sub> = 85	618	EC <sub>40</sub> = 425	EC <sub>30</sub> = 190	EC <sub>25</sub> = 32
3	Ag-Fe <sub>3</sub> O <sub>4</sub> NPs=0.5 g/l	618	EC <sub>40</sub> = 485	EC <sub>25</sub> = 370	EC <sub>15</sub> = 190	618	EC <sub>35</sub> = 375	EC <sub>20</sub> = 23	EC <sub>10</sub> = 135
	Ag-Fe <sub>3</sub> O <sub>4</sub> NPs=1 g/l	618	EC <sub>35</sub> = 325	EC <sub>20</sub> = 170	EC <sub>10</sub> = 70	618	EC <sub>30</sub> = 360	EC <sub>15</sub> = 140	EC <sub>5</sub> = 39
	Ag-Fe <sub>3</sub> O <sub>4</sub> NPs=1.5 g/l	618	EC <sub>45</sub> = 330	EC <sub>30</sub> = 170	EC <sub>20</sub> = 6.5	618	EC <sub>40</sub> = 400	EC <sub>25</sub> = 125	EC <sub>15</sub> = 33

\* EC values were calculated based on COD (mg/l).

#### 4.7.3. Direct Effects of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs Concentrations on the Acute Toxicity of Microtox and *Daphnia magna* in Pharmaceutical ww

The acute toxicity test was performed in the samples containing 0.5 g/l, 1 and 1.5 g/l Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentrations. In order to detect the direct responses of Microtox and *Daphnia magna* to the increasing Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentrations the toxicity test were performed without Pharmaceutical ww. The initial EC values and the the EC<sub>50</sub> values were measured in the samples containing increasing Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentrations after 150 min. Table 5 showed the responses of Microtox and *Daphnia magna* to increasing Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentrations.

The acute toxicity originating only from 0.5 g/l, 1 and 1.5 g/l Ag-

Fe<sub>3</sub>O<sub>4</sub> NPs concentrations were found to be low (Table 5). 0.5 g/l Ag-Fe<sub>3</sub>O<sub>4</sub> NPs did not exhibited toxicity to *Vibrio Fischeri* and *Daphnia magna* before and after 150 min. The toxicity attributed to the 1 g/l and 1.5 g/l Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentrations were found to be low in the samples without Pharmaceutical ww for the test organisms mentioned above. The acute toxicity originated from the Ag-Fe<sub>3</sub>O<sub>4</sub> NPs decreased significantly to EC<sub>1</sub> and EC<sub>4</sub> after 150 min. Therefore it can be concluded that the toxicity originating from the Ag-Fe<sub>3</sub>O<sub>4</sub> NPs is not significant and the real acute toxicity throughout the removal of microorganisms and organic compounds was attributed to the Pharmaceutical ww, to the microorganisms and organic compounds in Pharmaceutical ww, to their metabolites and to the disruption by-products (Table 5).

**Table 5:** The responses of Microtox and *Daphnia magna* acute toxicity tests in addition of increasing Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentrations without Pharmaceutical ww after 150 min.

Ag-Fe <sub>3</sub> O <sub>4</sub> NPs Conc. (g/l)	Microtox Test			<i>Daphnia magna</i> Test		
	Initial Acute Toxicity EC <sub>50</sub> Value (mg/l)	Inhibitions after 150 min	EC Values (mg/l)	Initial Acute Toxicity EC <sub>50</sub> Value (mg/l)	Inhibitions after 150 min	EC Values (mg/l)
0.5	EC <sub>10</sub> =25	-	-	EC <sub>10</sub> =30	-	-
1	EC <sub>15</sub> =130	2	EC <sub>1</sub> =5.00	EC <sub>20</sub> =150	5	EC <sub>3</sub> =8.00
1.5	EC <sub>20</sub> =210	5	EC <sub>4</sub> =10.00	EC <sub>30</sub> =250	7	EC <sub>7</sub> =12.00

## 5. Conclusions

This present study, the proposes a novel simple method was applied for the adsorption of ibuprofene and oxytetracycline in water using Ag-Fe<sub>3</sub>O<sub>4</sub> NPs. The TEM image of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs appeared darker than Fe<sub>3</sub>O<sub>4</sub> NPs. Our method of NPs synthesis could not produce uniform particles in size. The XRD patterns of Ag-Fe<sub>3</sub>O<sub>4</sub> NPs showed the 2θ values of 38.11°, 44.32°, 64.24°, 77.61° and 81.57° corresponding to the (111), (200), (220), (311) and (222) planes of cubic Ag, respectively. The Ms values was 61 emu/g for Fe<sub>3</sub>O<sub>4</sub> NPs and 69 emu/g for Ag-Fe<sub>3</sub>O<sub>4</sub> NPs. The increasing Ms values in the nanocomposite might have been due to the interactions between the NPs changed the anisotropic energy.

99% maximum ibuprofene adsorption removal yield was obtained at 2.03 mg/g maximum ibuprofene adsorption concentration, at pH = 7.0 and at 21°C, respectively. 99% maximum oxytetracycline adsorption removal yield was observed at 3.04 mg/g maximum oxytetracycline adsorption concentration, at pH = 5.0 and at 21°C, respectively. 93% maximum Ag-Fe<sub>3</sub>O<sub>4</sub> NPs adsorption yield was measured at 45 min contact time, 7 mg in 500 μl of suspension, at 0.1 mg/l Ag-Fe<sub>3</sub>O<sub>4</sub> NPs concentration at 3.0 mg/l ibuprofene and at 3.0 mg/l oxytetracycline, respectively. The characterization of the adsorbent by means of microscopy, spectroscopy and calorimetry techniques reveal the presence of Ag in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs and the adsorption of ibuprofene and oxytetracycline. The adsorption equilibrium is characterized by a Langmuir isotherm model.

94.44% maximum Microtox acute toxicity yield was observed in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1 g/l after 150 min, at 60°C. However, 90% maximum *Daphnia magna* acute toxicity yield was observed in Ag-Fe<sub>3</sub>O<sub>4</sub> NPs=1 g/l after 150 min, at 60°C. As a result, Microtox acute toxicity test yield is higher than *Daphnia magna* acute toxicity test for the removal of microorganisms and organic compounds from a Pharmaceutical ww using Ag-Fe<sub>3</sub>O<sub>4</sub> NPs. Therefore, it can be concluded that the toxicity originating from the Ag-Fe<sub>3</sub>O<sub>4</sub> NPs is not significant and the real acute toxicity throughout the removal of microorganisms and organic compounds was attributed to the Pharmaceutical ww, to the microorganisms and organic compounds in Pharmaceutical ww, to their metabolites and to the disruption by-products. Finally, the removal of microorganisms and organic compounds from a Pharmaceutical ww using Ag-Fe<sub>3</sub>O<sub>4</sub> NPs and the evaluation of Microtox and *Daphnia Magna* acute toxicity assays for these conditions was successfully implemented.

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