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Increasing the Efficiency of Magnetic Heterogeneous Fenton Catalysts with a Simple Halogen Visible Lamp

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Abstract:

The effect of a simple visible halogen lamp was studied in the Fenton-type oxidation of three model aqueous pollutants differing in their structure and electrostatic charge, methyloange (MO), methylene blue (MB) and paranitrophenol (PNP), using maghemite nanoparticles (γ -Fe₂O₃ NP), or maghemite/silica nanocomposite microspheres (γ -Fe₂O₃/SiO₂ MS) as heterogeneous catalysts. These materials, which were fully characterized, differ in size, morphology, porosity and microstructure, although their catalytic activity is related to the same γ -Fe₂O₃ nanoparticles. Both have a strong magnetic susceptibility, but only the MS catalyst can be easily recovered by magnetic settlement. Whatever the catalyst, the pollutant tested, or the experimental conditions used, much better decolorization rates and mineralization efficiencies were recorded under illumination by visible light in comparison to the same tests in the dark. The large range of experimental conditions tested enabled us to propose a mechanism for photocatalytic activity. Experiments of long-term stability showed that the MS catalyst, although generally less active than the NP catalyst, retained almost all of its activity after five repeated experiments under visible light. The good stability of this catalyst was also confirmed by the low level of iron leaching, making it suitable candidate for an application as photo-Fenton catalyst in industrial wastewater treatment.

Keywords:

Heterogeneous photo-Fenton, Iron oxide nanoparticles, Magnetic nanocomposite microspheres, Visible light irradiation, Wastewater.

1. Introduction

Nowadays, advanced oxidation processes (AOPs) are becoming increasingly important for wastewater treatment technologies, especially for poorly biodegradable contaminants.[1-5] The common feature of these processes is the production of extremely reactive and unselective radical oxygen species, especially the highly oxidizing hydroxyl radicals (HO^\bullet), which are able to degrade even the most recalcitrant molecules into biodegradable compounds, or mineralize them into CO_2 , H_2O and inorganic ions.[3] AOPs can be divided into different groups based on how the radicals are produced. One may mention the sonochemical processes, the radiolysis, the electrochemical methods, and the purely chemical processes which include the Fenton-type reactions ($\text{H}_2\text{O}_2/\text{Fe}$ catalyst). Nevertheless, one of the most important classes of AOPs is constituted by the photochemical processes, such as water, H_2O_2 and/or O_3 photolysis ($h\nu/\text{H}_2\text{O}$, $h\nu/\text{H}_2\text{O}_2$, $h\nu/\text{O}_3$, $h\nu/\text{H}_2\text{O}_2/\text{O}_3$), heterogeneous photocatalysis ($h\nu/\text{TiO}_2$ catalyst), and the photo-Fenton process ($h\nu/\text{H}_2\text{O}_2/\text{Fe}$ catalyst).[5,6] Their generally high efficiency is due to the assistance of UV or visible light irradiation, which is directly or indirectly responsible for the production of the radicals. UV light is usually used in the photochemical processes, because of its large energy which is able to induce the direct photolysis of an oxidant (H_2O_2 , O_3 , O_2 , $\text{H}_2\text{O}\dots$), or to activate a semi-conducting catalyst such as TiO_2 , leading to the generation of large amount of oxygen radicals. However, the photodegradation under UV irradiation is limited for industrial utilization, since natural UV occupies only 3–5% of the solar light energy that reaches earth, while artificial UV light generators consume large quantities of electrical power and can be expensive and unstable. Recently, much effort has been done to expand the response of the photodegradation processes to visible light irradiation, in replacement or addition to UV-light. Nevertheless, water-treatment by direct visible-light photolysis of an oxidant is unfavourable, since the rate of photolytic decomposition of the oxidants to oxygen radicals is very slow. To overcome this drawback, it is thus necessary to add a photocatalyst. [7] In recent years, there has been an extensive interest in the use of semiconductors in photochemical degradation. Among them, TiO_2 has been widely used as an effective photocatalyst for environment purification since it can decompose a large number of pollutants, [6-10] but its wide bandgap ($\sim 3.2\text{eV}$) limits its application to UV light irradiation. To the contrary, iron oxide particles (Fe_3O_4 , Fe_2O_3 , $\text{FeOOH}\dots$) are suitable candidates to be used as visible-light photocatalysts for water treatment. Indeed iron oxides have a relatively small bandgap ($\sim 2.1\text{ eV}$ for Fe_2O_3) which is appropriate for receiving visible light, and the electron-hole pair may generate

radical oxygen species in the same way as TiO_2 . [11-13] In addition, compared with TiO_2 , iron oxide presents the advantages to be non-toxic and inexpensive. [14,15] However it has been reported that iron oxides are relatively inefficient visible-light photocatalysts in the absence of additional oxidant, [11-13] while they are much more active as heterogeneous photo-Fenton catalysts, *i. e.* when H_2O_2 is added as additional oxidant to generate the oxygen reactive species. [13,16] To explain this result, it has been proposed that large amounts of HO^\bullet , are generated from H_2O_2 at the surface of the iron oxide in the presence of light. [17-19]

Although the photo-Fenton process is also efficient in homogeneous phase at $\text{pH}=3$ (*i. e.* with dissolved Fe^{2+} or Fe^{3+} as catalyst), the use of iron oxides as heterogeneous catalyst provides the possibility to recover and reuse the catalyst and to operate in a broader pH range. [20] In spite of these advantages, there is a relatively small number of studies on the use of iron oxides as photo-Fenton catalysts under visible light. [13, 17-29] These studies have clearly shown that the efficiency of the iron oxide/ H_2O_2 system in the degradation of various model pollutants is considerably increased through visible light irradiation, which might be slightly more efficient than UV light irradiation. [27] However the impact of the experimental conditions on the efficiency of visible light irradiation is still poorly understood. This article aims to highlight the parameters affecting the efficiency of visible light irradiation in a heterogeneous photo-Fenton process, using maghemite ($\gamma\text{-Fe}_2\text{O}_3$) nanoparticles (NP) as photocatalyst. The effect of the visible light irradiation on the activity of this catalyst is studied for a large range of conditions obtained by varying several parameters such as the presence of a silica support, the nature of the model pollutant, the pH, the H_2O_2 concentration, and the amount of catalyst. The choice of $\gamma\text{-Fe}_2\text{O}_3$ nanoparticles has been dictated by several considerations. First, $\gamma\text{-Fe}_2\text{O}_3$ is characterized by a high magnetic susceptibility, which offers an additional advantage in terms of recovery and reusability of the catalyst. Indeed magnetic iron oxides can be easily separated and removed from solution by simply applying an external magnetic field with a magnet or an electromagnet. [31-33] Magnetic settlement by an appropriate magnetic field provides a convenient and low cost method for the separation of solid particles in a suspension, as it is well illustrated in sewage or drinking water plants with the Sirofloc® process. [34] Second, it has been shown that $\gamma\text{-Fe}_2\text{O}_3$ and Fe_3O_4 are the most active and stable iron oxide phases in photo-Fenton processes. [16] Compared with Fe_3O_4 , $\gamma\text{-Fe}_2\text{O}_3$ nanoparticles are not submitted to oxidation, and are therefore more stable in water and air. Third, the $\gamma\text{-Fe}_2\text{O}_3$ nanoparticles used in this work have been thoroughly characterized and tested as Fenton catalyst in a previous publication. [35]

Among the parameters which are studied here, the presence of silica (SiO₂) microspheres (MS) as a porous support for the γ -Fe₂O₃ nanoparticles is of primary importance. It has been demonstrated that the dispersion of the catalyst on a support may considerably improve its activity, due to an increase of the adsorption of the pollutant and/or to a decrease of the recombination of the created electron-hole pairs.[28] Moreover, catalyst leaching problem can be avoided by hosting the heterogeneous catalyst onto a support.[36] In the case of a magnetic catalyst, we have shown in addition that the presence of a support can favor the recovery of the catalyst by magnetic settlement.[35] To study how the efficiency of light irradiation is affected by the nature of the pollutant, we chose to compare the degradation of three aqueous model pollutants, methylene blue (MB), methyl orange (MO) and paranitrophenol (PNP), which can be distinguished in terms of their charge, their light absorption properties, and their chemical structure. Finally, a simple halogen lamp has been taken as source of visible light. We show that this simple and cheap lamp is able to considerably increase both the decolorization and mineralization rate of the pollutant solutions without detrimental impact on the stability of the heterogeneous catalyst.

2. Experimental

2.1 Catalysts synthesis

The synthesis of the catalysts has been already described in our previous paper. [35] Briefly, magnetite (Fe₃O₄) NP were first prepared by the Massart's method, adding ammonia to an aqueous mixture of FeCl₃ and FeCl₂. [37] Then, the Fe₃O₄ NP were oxidized to γ -Fe₂O₃ by successively adding HNO₃ and Fe(NO₃)₃. [38-39] The γ -Fe₂O₃ NP were finally obtained as an aqueous acidic dispersion with a high colloidal stability, also called ferrofluid. The γ -Fe₂O₃ NP were supported on the surface of porous silica microspheres via a protocol inspired from the method originally described by Andersson *et al.* [35,40] Briefly, a magnetic sol containing the γ -Fe₂O₃ NP and a silica precursor (Tetraethoxysilane, TEOS) in acidic medium was added dropwise to an organic phase composed of a vegetable oil (usually commercial rapeseed oil) and a commercial emulsifier (Arlacel P135 from Uniquema), under stirring. The water-in-oil emulsion thus formed was transferred into a Büchner flask, and the ethanol formed by the hydrolysis of TEOS was evaporated from the dispersed phase under reduced pressure, which led to a rapid condensation of TEOS in silica in each water droplet. The emulsion was then broken by addition of a large amount of acetone. The silica MS containing the γ -Fe₂O₃ NP were washed several times with acetone and water, and dried at 70 °C. Finally, the beads were

calcined in an oven under air at 400 °C for 18 h to remove any organic traces. In the following sections, the unsupported γ -Fe₂O₃ nanoparticles will be referred as the γ -Fe₂O₃NP catalyst, while the γ -Fe₂O₃ nanoparticles supported on the surface of the silica microspheres will be referred as the γ -Fe₂O₃/SiO₂ MS catalyst.

2.2 Characterization of the catalysts

Transmission electron microscopy (TEM) micrographs of the γ -Fe₂O₃NP and γ -Fe₂O₃/SiO₂ MS catalysts were taken using a JEM JEOL 100 CX microscope operating at 100 kV. In this aim, the MS were embedded in a resin (AGAR 100), which was then polymerised at 60°C during two days, and cut in 70 nm thin sections using a LEICA ULTRACUT UCT microtome apparatus. The MS catalysts were also observed by scanning electron microscopy (SEM) on a SEM-FEG Hitachi SU-70 apparatus. The images were taken in secondary electron mode with an accelerating voltage of 10 kV. Prior to analysis, the beads were coated with a thin shell of gold by sputter deposition. The size distribution of the particles was determined from TEM or SEM pictures analyzed using the Image J software. The amount of γ -Fe₂O₃NP in the MS was determined by spectrophotometry. In this aim, a given weight of material was first introduced in an HCl aqueous solution (4 mol L⁻¹). After 48 h of stirring, the γ -Fe₂O₃ NP were totally dissolved, and the iron concentration in the supernatant was analyzed by atomic absorption spectrophotometry (Perkin Elmer AA100 apparatus) and the weight fraction of γ -Fe₂O₃ in the beads, $w_{\text{Fe}_2\text{O}_3}$ was determined. A similar procedure was used to determine the amount of γ -Fe₂O₃NP dispersed in water. Magnetic characterizations of the NP and MS catalysts were respectively made at 25 °C on a vibrating sample magnetometer (home-made apparatus) and SQUID magnetometer (Quantum Design MPM-5S apparatus). The NP were analysed dispersed in water, whereas the MS were analysed in powdered form. XRD spectra of the powdered samples were recorded using a Phillips PW 1130 diffractometer. Data were collected from $2\theta = 10^\circ$ to 80° in 0.1° steps. The room-temperature UV-visible-NIR spectra of the powdered samples were recorded with a diffuse reflectance cell (internal sphere) in the range 200-2500 nm on a Varian Cary 400 spectrometer (Teflon as a blank). The pure γ -Fe₂O₃ NP were dispersed in Teflon before analysis. Volumetric adsorptions of nitrogen at 77 K and CO₂ at 273 K were performed on a ASAP 2020 Micromeritics apparatus. Before analysis, the samples were degassed overnight at 110°C under high vacuum (10^{-6} bar).

2.3 Photocatalytic activity tests

The photoactivity of the two catalysts ($\gamma\text{-Fe}_2\text{O}_3$ NP and $\gamma\text{-Fe}_2\text{O}_3/\text{SiO}_2$ MS) was tested for the degradation of three aqueous pollutants, methylene blue (MB), methyl orange (MO), or paranitrophenol (PNP). The chemical structure of these three compounds can be found in the ESI (fig. S1). All of the experimental investigations were carried out in a cylindrical double jacketed glass reactor (4 cm diameter, 6 cm height) with 20 mL of reaction volume in contact with the atmospheric air (see the photograph and scheme of the reactor in fig. S2 of the ESI). The reactor was maintained at a fixed temperature ($T = 40^\circ\text{C}$), which was controlled by circulating water. This temperature was chosen due to several studies which have reported that the temperature range $35\text{-}45^\circ\text{C}$ is an optimum to promote the degradation and mineralization of pollutants in Fenton-type processes.[41] Below these temperatures, the rate of production of HO^\bullet radicals is limited, which decreases the removal efficiency of the catalyst, whereas above these temperatures, it has been found that the catalytic activity can be decreased, due to the increase of decomposition rate of H_2O_2 in oxygen and water without producing HO^\bullet radicals. We also performed some preliminary Fenton tests on the MS catalyst, using MB as model pollutant. These tests have also shown that the performance of the MS catalyst is lower at $T=30^\circ\text{C}$ than at $T=40^\circ\text{C}$.

The visible irradiation source was an Osram 46W halogen lamp vertically placed in the center of the reactor, and surrounded by the two glass jackets and the circulating water. The visible-light illuminance in the center of the photoreactor (75 ± 3 Klux) was measured with a digital TES 1335 light meter. The UV-visible emission spectrum of the halogen lamp was taken using an Ocean Optics S2000 spectrometer (for the spectrum, see fig. S3 in the ESI). The relative spectral irradiance at each wavelength (raw measurements) was converted into absolute spectral irradiance through the use of the Abney equation (see the appendix in the ESI). [42] The integration of the spectrum from $\lambda=200$ nm to 880 nm enables to estimate the total irradiance (in the UV, visible and NIR part of the spectrum), which is equal to $E_e = 480$ W m^{-2} . While a halogen lamp can emit a small amount of UV light, the emission spectrum of the lamp showed that the UV light intensity is negligible in the photoreactor, which can be explained by the natural filter formed by the double glass jackets and the circulating water.

The standard photocatalytic tests were conducted as follows. 0.6g of $\gamma\text{-Fe}_2\text{O}_3/\text{SiO}_2$ MS, or 420 μL of the $\gamma\text{-Fe}_2\text{O}_3$ NP acidic aqueous dispersion (both equivalent to $[\text{Fe}]_T=3.15\times 10^{-2}$ mol L^{-1} in the final reaction mixture) was added to an aqueous solution of the pollutant, which pH was previously adjusted at 3 by addition of HNO_3 . The volume of this solution was chosen in a

manner that the initial concentration of the pollutant in the final mixture was $C_i = 2.5 \times 10^{-4}$ mol L⁻¹ for a total volume of 20 mL (except for the MB pollutant with the MS catalyst, where C_i was fixed at 8×10^{-4} mol L⁻¹ to take into account the much larger adsorption of this compound on the silica surface). Before turning on the lamp, the suspension was magnetically stirred for 2h to establish the adsorption equilibrium. Then the photocatalytic oxidation was initiated by turning on the halogen lamp and by simultaneously adding H₂O₂ (1.36 mL of a 30%, (w/w) commercial aqueous solution, corresponding to an initial concentration of H₂O₂ of 0.68 mol L⁻¹) to the system. The solution samples were taken at desired time intervals, and were put over a magnet to separate the supernatant from the catalysts by magnetic settlement. For the γ -Fe₂O₃ NP, a concentrated KCl aqueous solution (50 g L⁻¹) was previously added to make aggregation of the nanoparticles and thus to favour their settlement. The supernatant was then recovered and diluted to an adequate concentration before to be analyzed with an UV-visible spectrophotometer (UVIKONXL apparatus). The remaining concentration of the pollutant in the supernatant (C_t) was determined using the Beer–Lambert law at 502, 400, and 665 nm respectively for MO, PNP, and MB. Time zero for reaction take-off was defined as the time when the lamp was turned-on and H₂O₂ simultaneously added. The concentration of the pollutant in the supernatant measured at t=0 was noted C_0 . C_0 varied from 1.8×10^{-4} to 2.5×10^{-4} mol L⁻¹, depending on the amount of adsorbed pollutant on the catalyst. The initial rate of decolorization (v_0) was determined by plotting the tangent at t=0 of the kinetic curve $C_t = f(t)$. The decolorization yield (DY) at 3h, was also evaluated from this curve, using the following equation (1).

$$DY = 100 - 100 \cdot C_{3h} / C_0 \quad (\text{Eq. 1})$$

Where C_{3h} and C_0 are the concentrations of pollutant in the supernatant respectively at t=3h and t=0. Since most of the pollutant should ultimately break down into inorganic ions and molecules, an important aspect of this study is also to evaluate the mineralization of the system. The mineralization yield of the pollutant at 24h (MY) was monitored by analyzing the non-purgeable organic carbon (NPOC) in the supernatant using a Shimadzu TOCASI-5000A apparatus and was calculated using the equation (2).

$$MY = 100 - 100 \cdot \text{NPOC}_{24h} / \text{NPOC}_i \quad (\text{Eq. 2})$$

Where NPOC_{24h} is the NPOC concentration (in ppm) in the supernatant at t = 24 h, and NPOC_i the initial NPOC concentration (before the adsorption equilibrium). The amount of iron ions leached from the catalysts in the solution was determined by measuring the iron

concentrations in the supernatant at 4h using atomic absorption spectrophotometry. The percentage of leached iron (Fe_L) was calculated using the following equation (3).

$$Fe_L = 100 \cdot [Fe]_{s,4h} / [Fe]_T \quad (\text{Eq. 3})$$

Where $[Fe]_{s,4h}$ is the iron concentration in the supernatant at 4h, and $[Fe]_T$ the equivalent iron concentration used in the catalytic test ($[Fe]_T = 3.15 \times 10^{-2} \text{ mol L}^{-1}$).

Some control experiments in dark were also carried out to study the performance of the Fenton process on the oxidation of the three pollutants, and to evaluate the effect of visible light on the performance of two catalysts. The catalytic and photocatalytic tests at pH=5 and 8 were performed by replacing the HNO_3 solution by pure water and NaOH solution, respectively. The reusability of the $\gamma\text{-Fe}_2\text{O}_3/\text{SiO}_2$ MS catalyst for four times was evaluated using MO as model pollutant, in absence and in presence of light. The reuse tests were performed as follow. After each catalytic test (corresponding to 4 hours of reaction), the catalyst was separated from the solution by magnetic settlement, rinsed two times with $10^{-3} \text{ mol L}^{-1} HNO_3$ aqueous solution and two times with water to remove any possible contaminant from the surface. The solid was finally dried in an oven at 70°C overnight before the next use. Some additional tests were carried out to measure the activity of the catalysts on the decomposition rate of H_2O_2 in absence of any pollutant. The same experimental conditions were used, except that no pollutant solution was added. The solution samples were taken at desired time intervals, and the catalyst was separated from the supernatant by the same procedure. Then the H_2O_2 concentration in the supernatant was determined by the following procedure. 0.8 mL of the supernatant was transferred in a 10 mL graduated flask containing a large amount of water and 0.28 mL of H_2SO_4 98% solution. The total volume was completed by adding water, in such way that the dilution factor for H_2O_2 was 12.5, and the concentration of H_2SO_4 was 0.5 mol L^{-1} . An aliquot of this solution was then quickly titrated by a $0.02 \text{ mol L}^{-1} KMnO_4$ solution. At the equivalent point, which is detected by the persistence of the pink coloration due to $KMnO_4$, the stoichiometric factor between the amounts of H_2O_2 and $KMnO_4$ is 2.5.

3. Results and discussion

3.1 Characterization of the catalyst

The characterization of the catalysts has been thoroughly described in our previous paper.[35] The main results of this study are summarized in the following paragraph. The morphology of

the catalysts was examined by SEM and TEM, as depicted in figure 1 (A and B). Fig.1A shows that the γ -Fe₂O₃NP are polydisperse in size and have a rock-like morphology, which can be approximated as spheres having a mean diameter of $d=8.9$ nm with a standard deviation of 2.4 nm, as obtained from the digitized TEM image of 170 particles in combination with an image analysis software. SEM image of the γ -Fe₂O₃/SiO₂ MS (fig. 1B), shows that the silica microsphere supports are polydisperse in size with a relatively smooth external surface. Their mean diameter determined by counting the size of more than 350 beads is $D=2.0$ μ m with a standard deviation of 1.7 μ m. The TEM image of a bead (see the inset on fig. 1B) shows a homogeneous and isotropic dispersion of the γ -Fe₂O₃NP in the silica, without evidence for the formation of agglomerates or chains, despite the relative high weight fraction of the γ -Fe₂O₃ NP in the microspheres (a value of $w_{\text{Fe}_2\text{O}_3}=8.1\%$ w/w is found by atomic absorption spectrophotometry).

Magnetic properties of bare γ -Fe₂O₃NP and γ -Fe₂O₃/SiO₂ MS were respectively investigated with a vibrating sample magnetometer (VSM) and SQUID magnetometer. The absence of hysteresis in the magnetization curves shows that the γ -Fe₂O₃ nanoparticles exhibit typical superparamagnetic behaviour (see the curves of fig.S4-A in the ESI), both in the absence and the presence of the silica support. This proves that the magnetic properties of the nanoparticles were preserved after their encapsulation in the silica microspheres. Both curves show in addition a strong magnetic susceptibility. This strong magnetic susceptibility is spectacularly confirmed by the behaviour of the samples in water, in presence of a magnet (see the photographs in fig.1C and 1D). Both materials are strongly attracted by the magnetic force proportional to the strong magnetic field gradient. However the very high colloidal stability of the NP in water restricts their separation from the water, on the contrary of the MS particles which can be completely separated from the solution in a very short time (less than 3 min). The magnetic saturation values determined from the curves are 62 and 66 emu g⁻¹, respectively for the NP and the MS samples (the magnetic saturations are normalised to 1 g of maghemite). Both values agree well with the value usually given for γ -Fe₂O₃ nanoparticles ($M_s= 60$ emu g⁻¹). [43] The diffraction patterns (see fig. S4-B) obtained for the magnetic nanoparticles before and after their supporting in the silica microspheres are similar, and corresponds well to the spinel structure of maghemite, confirming that the γ -Fe₂O₃ nanoparticles are not altered by their encapsulation in the silica matrix. The average crystal size of nanoparticles calculated from the XRD pattern using the Scherrer formula is $d=7.1$ nm, which agrees well with the diameter of the γ -Fe₂O₃ NP obtained by TEM image. The absence

of variety of iron other than maghemite is also confirmed by the UV-Vis-NIR spectra of the γ - Fe_2O_3 NP and γ - $\text{Fe}_2\text{O}_3/\text{SiO}_2$ MS catalysts (see fig.S4-C).

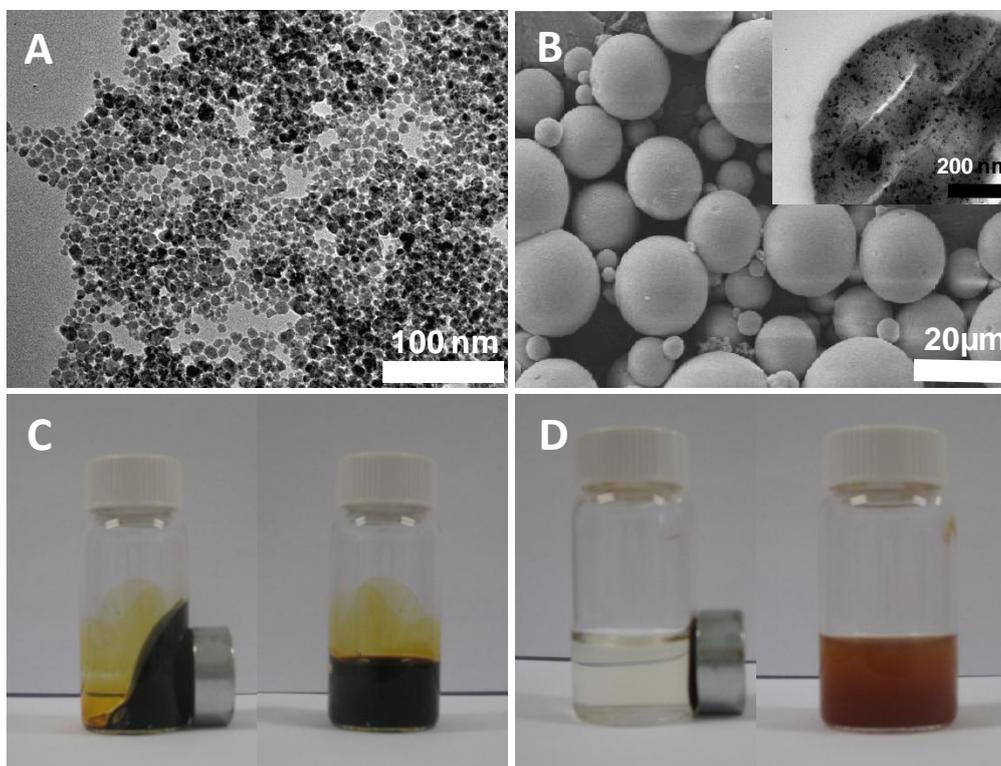


Fig.1: (A) TEM image of the γ - Fe_2O_3 NP. (B) SEM image of the γ - $\text{Fe}_2\text{O}_3/\text{SiO}_2$ MS. In inset: TEM image of a γ - $\text{Fe}_2\text{O}_3/\text{SiO}_2$ MS. (C) Photograph of the water dispersion of the γ - Fe_2O_3 NP (with/ without magnet). (D) Photograph of the water suspension of the γ - $\text{Fe}_2\text{O}_3/\text{SiO}_2$ MS (with/without magnet).

The porosity of the materials was determined by sorption volumetry of N_2 and CO_2 (respectively at 77 K and 273 K). Both are characterized by a large surface area, which is of prime importance to obtain some high catalytic activities. For the γ - Fe_2O_3 NP, the N_2 isotherm (see fig. S4-D) is of type II, which is observed for finely divided non-porous solids. By applying the BET equation to the first part of the isotherm, it was found that the specific surface area is $159 \text{ m}^2 \text{ g}^{-1}$, which corresponds to the theoretical external surface of spherical particles having a mean diameter of $d=7.4 \text{ nm}$. For the γ - $\text{Fe}_2\text{O}_3/\text{SiO}_2$ MS, the N_2 isotherm (see fig. S4-D) is of type I, which is characteristic of a microporous solid with pore diameters smaller than 2 nm. The calculus of the BET surface area from the N_2 isotherm being inoperative for microporous solids, CO_2 adsorption volumetry was practiced on the MS.[35] By modelling this isotherm with the Dubinin-Astakhov equation,[44] a surface area (S) of 744

$\text{m}^2 \text{g}^{-1}$ and a microporous volume (V_p) of 0.30 mL g^{-1} were found for the beads. These values confirm the large amount of micropores in the silica network. In addition, the mean diameter of the micropores, D_p , was calculated using the relation $D_p = 4V_p/S$. A value of $D_p = 1.6 \text{ nm}$ was found, showing that the micropores have a relatively large size, which may allow the diffusion of the pollutants into the beads.

3.2 Photocatalytic tests on the three pollutants

The performance of the two catalysts, $\gamma\text{-Fe}_2\text{O}_3$ NP, and $\gamma\text{-Fe}_2\text{O}_3/\text{SiO}_2$ MS, was examined under both dark (heterogeneous Fenton process) and visible light (heterogeneous photo-Fenton process) conditions. Their activity on the oxidation by H_2O_2 of the three aqueous pollutants (MO, MB, and PNP) was first tested at $\text{pH}=3$ and $T=40^\circ\text{C}$. H_2O_2 was added 2 h after the adsorption of the pollutant on the catalyst ($t=0$). The decolorization and the mineralization kinetics of the solutions were respectively followed by UV-visible spectroscopy, and measurement of the NPOC. The iron leaching of the catalysts was measured by analysis of the iron concentration in the supernatants after 4 h, using atomic absorption spectroscopy. In addition, to estimate the activity of the catalysts independently of the nature of the pollutant, the decomposition rate of H_2O_2 was determined by measurement of the H_2O_2 concentration at fixed times, using KMnO_4 titration, in absence of the pollutants. Figure 2 shows the kinetic curves obtained for the MO decolorization (fig.2A), the MO mineralization (fig.2B), and the H_2O_2 decomposition (fig.2C) in the supernatant. Other kinetic curves (MB and PNP decolorization and mineralization) and the resulting data can be found in the ESI (fig. S5 and table S1).

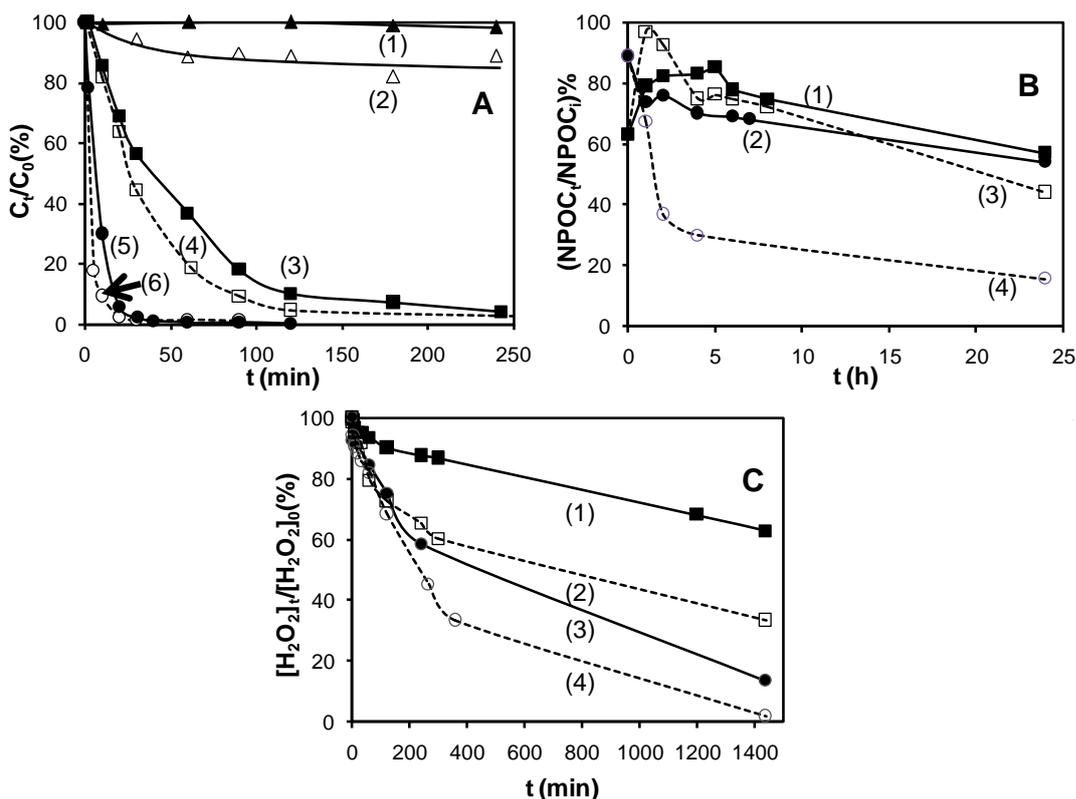


Fig.2 : (A) Kinetic curves for the decolorization of the MO solutions (The MO concentration, C_t is normalized to C_0 , the MO concentration at $t=0$) : (1) = without catalyst, absence of light, (2) = without catalyst, presence of the visible light, (3) = MS catalyst, Fenton, (4) = MS catalyst, photo-Fenton, (5) = NP catalyst, Fenton, (6) = NP catalyst, photo-Fenton. (B) Kinetic curves for the mineralization of the MO solutions (The NPOC concentration, $NPOC_t$ is normalized to $NPOC_i$, the initial NPOC concentration, before adsorption equilibrium) (1) MS catalyst, Fenton, (2) NP catalyst, Fenton, (3) MS catalyst, photo-Fenton, (4) NP catalyst, photo-Fenton. (C) Kinetic curves for the H_2O_2 decomposition (The H_2O_2 concentration, $[H_2O_2]_t$ is normalized to $[H_2O_2]_0$, the H_2O_2 concentration at $t=0$) : (1) = MS catalyst, Fenton, (2) = MS catalyst, photo-Fenton, (3) = NP catalyst, Fenton, (4) = NP catalyst, photo-Fenton.

In absence of catalyst, the MO solutions remain almost stable under the addition of H_2O_2 even in presence of illumination by visible light (see curves (1) and (2), fig. 2A). To the contrary, the addition of the MS or NP catalyst allows nearly total decolorizations of the MO solutions after few hours (see curves (3) to (6), fig.2A), and significant mineralization yields after 24h (see fig.2B), which is related to important decomposition rates of H_2O_2 (see fig.2C). To explain the correlation between the MO oxidation (fig.2A and 2B) and the H_2O_2

decomposition (fig. 2C) we have to take into account that a faster H_2O_2 decomposition leads to a larger production of HO^\bullet radicals which are responsible for the MO oxidation. The mineralization of the MO solutions (fig. 2B) occurs much more slowly than their decolorization (fig.2A), as already mentioned by other authors.[16,45] The best mineralization yields (55.9% for the MS catalyst and 84.5% for the NP catalyst) are obtained at 24h for the photo-Fenton process, clearly showing the interest of using a visible lamp. The patterns of the mineralization curves are rather complex. At $t=0$, the NPOC values are smaller than expected, because of the adsorption of a small part of the pollutant on the NP or MS catalysts. During the first hours of the reaction, the amount of NPOC is increased, especially for the MS catalyst, which can be interpreted in terms of desorption of the primary organic species resulting from the early degradation of the adsorbed MO. Finally, the NPOC is gradually decreased up to 24h, which is due to the slow mineralization of these primary organic species. However, the same trends are observed by the three measurements (decolorization, mineralization, and H_2O_2 decomposition). First, the NP catalyst is more efficient than the MS catalyst, in presence or in absence of light. The lower activity of the MS catalyst can be explained by the difficulty for the reactants to access to the surface of the $\gamma\text{-Fe}_2\text{O}_3$ NP, encapsulated in the core of the silica microspheres. Second, the efficiency of the two catalysts is greatly improved by irradiating the suspensions with visible light. This positive influence of the visible irradiation is due to an increase of the production of the HO^\bullet radicals at the surface of the catalyst, as already observed by several authors for similar processes.[18,22,25,26] Interestingly, this increase occurs without additional Fe leaching, which remains less than 0.4% (see table S1 in the ESI part). These results can be generalized to the two other pollutants, MB and PNP. Figure 3 shows the initial decolorization rates (v_0 , fig. 3A) and the mineralization yields at 24h (MY, fig. 3B) obtained for the three pollutants with the two catalysts, in the absence and in the presence of visible light.

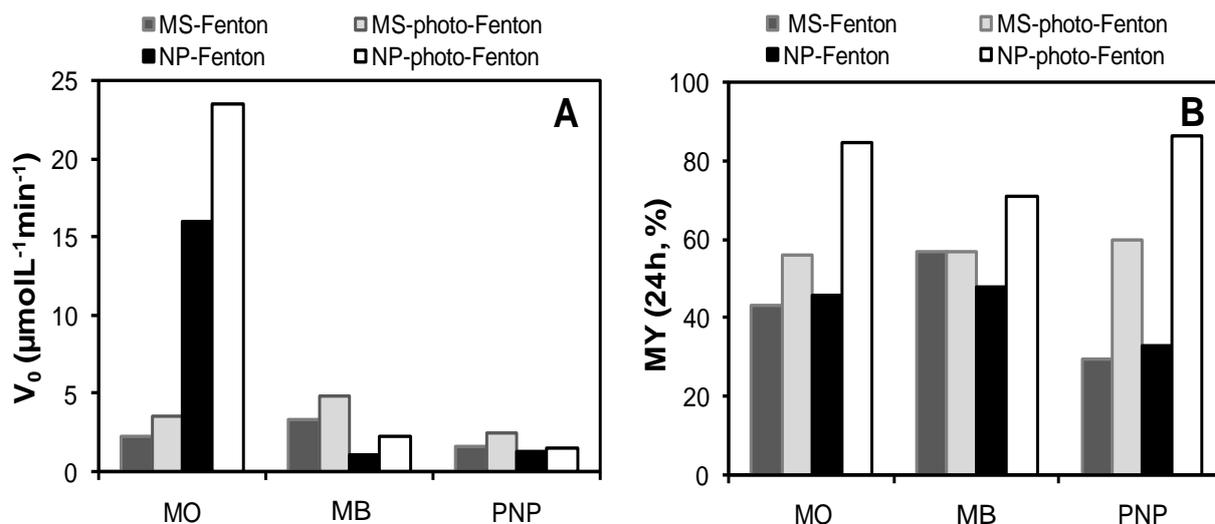


Fig.3 : (A) Initial decolorization rates (v_0) for the three pollutants (MO, MB, and PNP). (B) Mineralization yields (MY) at 24 h for the three pollutants (MO, MB, and PNP).

The decolorization and mineralization rates strongly vary with the pollutant, and it is difficult to highlight a clear trend, which depends on the catalyst, the process, and what it is measured (decolorization or mineralization). The NP catalyst is generally more efficient than the MS catalyst, but the results for PNP and MB are not as straightforward as for MO or for H_2O_2 decomposition, especially regarding decolorization. In some cases, comparable or even larger rates can be obtained with the MS catalyst, despite the fact that the accessibility of the reactants to the catalytic sites is lowered by the presence of the silica. This demonstrates that the decolorization rate is influenced by other parameters, such as the adsorption level of the pollutant, its stability to the attack of the HO^\bullet radicals, and the existence of several colored degradation intermediates at the first stages of the process. In a recent paper, it has been shown in particular that the reactivity of a pollutant toward an AOP can dramatically change, depending on the electrostatic interactions between the pollutant and the support. [46] The oxidation is generally favored when the charges are opposite. This can explain why in our case the decolorization of MB, a positively charged molecule, is increased in presence of silica, a negatively charged support. Nevertheless, all the data clearly show that the Fenton-like oxidation of the pollutants is considerably improved through the use of the visible lamp, although the importance of this improvement vary with the pollutant, the catalyst and the measured parameter. Concerning the decolorization rates (fig.3A), v_0 is about 1.2 to 2.1 times faster for the NP catalyst, and 1.5 to 1.6 times faster for the MS catalyst when visible lamp is

used. These higher rates lead to quasi-quantitative decolorization yields in only 3h, while it takes at least 4h for the reaction in the dark (see table S1). It is important to note that the level of v_0 increase due to visible light (fig. 3A) is comparable to the level of increase in H_2O_2 degradation rate (fig. 2C), which is of same order of magnitude (around 1.5 to 2). Therefore, during the early stages of the degradation, which corresponds to the decolorization, the efficiency of irradiation seems to be poorly influenced by the presence of the pollutant.

For mineralization (fig. 3B), best results are obtained for the NP catalyst, with very good mineralization yields under visible light (MY= 70.8% to 86.3%). For the MS catalyst, we also noted somewhat good mineralization yields in the same conditions (MY= 57.0% to 59.8%). The lower MY values obtained for the two catalysts in the absence of light (MY=29.3% to 56.9%) indicate that the dark Fenton process is not effective for the mineralization of the pollutants, a major limitation of this process, as also observed in relevant studies. [47,48]

Thus, the level of mineralization is generally well correlated to the decolorization rate : a high initial v_0 generally leads to a MY at 24H00 . However there is no simple relationship between the initial decolorization rate and the amount of mineralized pollutant, and sometimes a relatively moderate v_0 leads to relatively good MY at 24H00, as it is observed for example in the case of MB. This is explained by the great difference between the processes respectively measured by decolorization and mineralization. When measuring the decolorization, we measure in fact the result of few competitive reactions between the HO^\bullet radicals and the model pollutant (MO, MB or PNP), occurring during the few first hours and leading to uncolored intermediates having a chemical structure relatively close to the parent molecule. To the contrary, mineralization is the result of numerous competitive and consecutives oxidative reactions occurring during 24 hours, which lead finally to CO_2 , H_2O and inorganic salts. The kinetic of this complex process depends on the reactivity of all the formed intermediates towards the HO^\bullet radicals (and also towards other radical species such as HO_2^\bullet or O_2^\bullet). The nature of these intermediates depends on the nature of the parent molecule but also on the presence or absence of light, and on the nature of the catalyst. Therefore, in some cases, the uncolored intermediates can be formed relatively slowly due to the relative stability of the parent molecule towards oxidation, which lead to moderate v_0 , but they may react relatively fast with the HO^\bullet radicals, which leads to relatively high values of MY after 24H00.

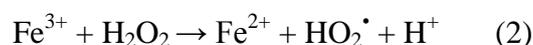
It should also be noted that the best MY gains due to visible light are always obtained with the NP catalysts, showing a better efficiency of the illumination for this catalyst. The reduced

efficiency of visible light with the MS catalyst may be caused by the larger turbidity of the solution, due to the large size of the γ -Fe₂O₃/SiO₂ particles, which may slightly decrease the penetration of visible light into the reactor. This is contradictory to what has been observed for decolorization, probably because this effect is appreciable only at long term, after several hours of illumination. Concerning the influence of the pollutant on the irradiation efficiency, larger gains are generally obtained for PNP, where mineralization yields are increased by a factor 2 to 2.5. To the contrary, the gain obtained for MB with the MS catalyst is negligible. These differences cannot be explained by the lower light absorption of PNP ($\epsilon L = 9531 \text{ L mol}^{-1}$ at $\lambda_{max} = 317 \text{ nm}$) in comparison to MO or MB ($\epsilon L = 34500 \text{ L mol}^{-1}$ and 79600 L mol^{-1} at $\lambda_{max} = 502 \text{ nm}$ and 665 nm , respectively), since mineralization mainly occurs later after decolorization. Here again, the explanation could be in the different mineralization pathways of the pollutants, which may vary in the presence of light. The difference could be especially due to the changes in the concentration of compounds recalcitrant to mineralization, such as oxalic, formic or acetic acid which have been identified as major end-products during Fenton and photo-Fenton processes. [49, 50]

From these results, a discussion can be made about the mechanism of the catalytic activity under visible light. Several studies have demonstrated that the higher efficiency of photo-Fenton process is due to the formation of more HO[•] radicals than in the dark Fenton process. In absence of light, the production of HO[•] is mainly due to the following reaction ($k_{obs} = 76 \text{ L mol}^{-1} \text{ s}^{-1}$ in homogenous phase). [51, 52]

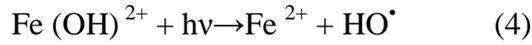


The catalytic cycle is completed by the following reaction:



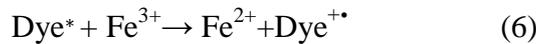
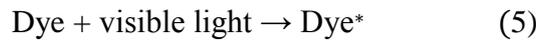
But this last step is relatively slow ($k_{obs} = 0.01 \text{ L mol}^{-1} \text{ s}^{-1}$, in homogenous phase) which decreases the efficiency of the iron based catalysts in the dark. In the presence of light, three pathways have been proposed to explain the increase of the catalytic activity. The first mechanism (reactions (3)-(4)) is due to the photo-reduction of Fe(III) to Fe(II) which accelerates the regeneration of Fe²⁺ ions, and at the same time produces additional HO[•] radicals. [53]



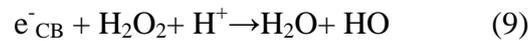
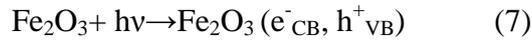


Since in homogenous phase, the $\text{Fe}(\text{OH})^{2+}$ mainly absorbs in the UV region, the second step (reaction (4)) is operative under UV irradiation ($\lambda \approx 310 \text{ nm}$).

In the second pathway (reactions (5)-(6)), called “dye-sensitization”, a colored molecule (a dye) can absorb visible light energy, thus being activated to an excited state. Then the excited dye molecule can transfer an electron to $\text{Fe}(\text{III})$ to generate $\text{Fe}(\text{II})$, which can promote the $\text{Fe}(\text{III})/\text{Fe}(\text{II})$ catalytic cycle.[54,55]



The third pathway (reactions (7)-(9)) is based on the semiconducting properties of iron oxides. When the band gap of Fe_2O_3 is illuminated by UV or visible light, electron-hole pairs are generated. Then the electrons present in the conduction band can be trapped by the surface Fe^{3+} leading to Fe^{2+} , or can react with H_2O_2 forming HO^{\bullet} radicals. [12, 48]



The first proposed mechanism is inoperative to explain our results, since photoreduction of the Fe^{3+} ions only occurs under UV light. The dye-sensitization mechanism is not predominant, at least in the beginning of the oxidative process. Indeed, during the early stages of the degradation of the pollutant - which are detected by the measurement of the decolorization - it seems that the increase of the catalytic activity induced by visible light is poorly influenced by the presence of the pollutant. In other words, the absorption of a part of the visible light by the pollutant has no impact on the photocatalytic activity. To the contrary, the fact that in the absence of any pollutant, visible light has a similar impact on the rate of H_2O_2 degradation stands for a mechanism principally based on photogenerated electron-hole pairs (reactions (7)-(9)). To explain that reaction (7) can be promoted by visible light, it has to be considered that the band gap of $\gamma\text{-Fe}_2\text{O}_3$ is sufficiently low ($E_g = 2.2 \text{ eV}$) to be sensitive to visible light. Thus, our results are in good agreement with a predominant mechanism based on photogenerated electron-hole pairs, although a more complex pathway associating photogenerated electron-hole pairs and dye-sensitization may be suggested to explain the

effect of visible light, and especially the differences in the mineralization yields which result from the late stages of the degradation process.

3.3 Influence of experimental parameters

Further studies were carried out to evaluate the effect of different reaction variables, especially the solution pH, the hydrogen peroxide concentration, the amount of catalyst and the light intensity of the lamp. We examined the influence of these parameters on the decolorization kinetics, using MO as model pollutant. The results concerning the influence of the pH, H_2O_2 concentration, and amount of catalyst on v_0 are given in figure 4. The results concerning the influence of light intensity can be found in ESI (figure S6).

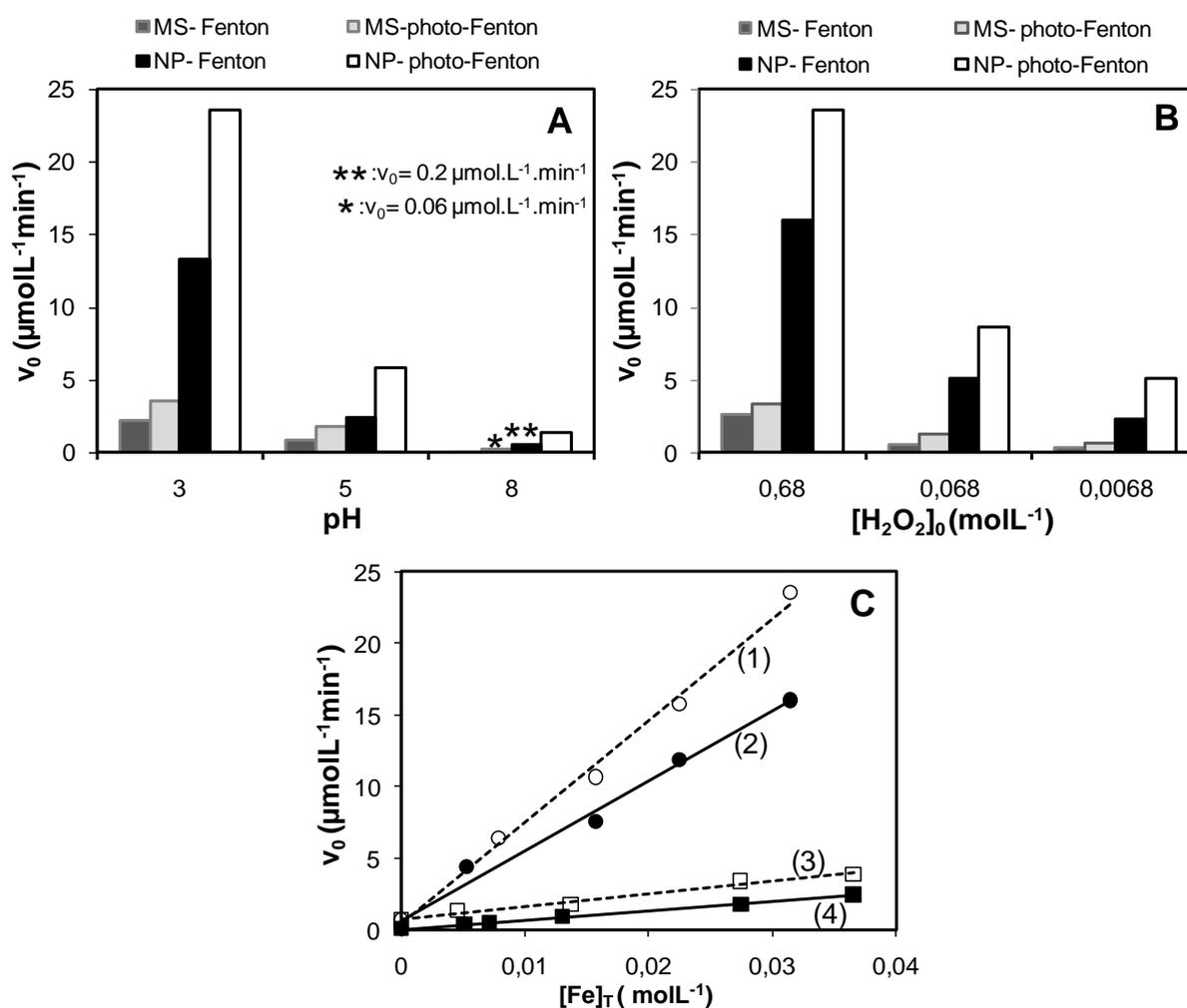


Fig.4: Influence of various parameters on the initial decolorization rate (v_0), using MO as model pollutant. (A) pH. (B) Initial concentration of H_2O_2 ($[H_2O_2]_0$). (C) Amount of catalyst ($[Fe]_T$) : (1) NP-photo-Fenton, (2) NP-Fenton, (3) MS-photo-Fenton, (4) MS-Fenton.

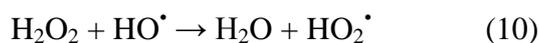
3.3.1 Effect of the pH

The pH is an important parameter to study, since the Fenton and photo-Fenton reactions are strongly pH dependent. It has been mentioned in the literature that the generation of the HO[•] radicals, and thus the efficiency of the oxidation is optimum at acidic pH and decreases as the pH increases. [20] The decrease of the catalytic activity in homogeneous phase when the pH is increased, is also explained by the precipitation of the soluble the Fe²⁺ and Fe³⁺ ions as poorly active iron (II) and (III) hydroxides. Therefore, the application of homogeneous Fenton catalysts is generally limited to a narrow pH range (around pH=3), and the heterogenization of this process is of prime importance to obtain a catalyst capable of maintaining its efficiency in a large pH range. Our results concerning the effect of the pH are shown in fig. 4A. This figure shows that the pH significantly influences the degradation of MO. It can be seen that the increase of the pH from 3 to 8 leads to a strong decrease of the decolorization rate, independently of the catalyst (MS or NP) or the presence of light. This result is in agreement with other works carried out on iron oxide based heterogeneous Fenton catalysts.[16,18,25,26,48] The highest decolorization rate is obtained at pH=3. When increasing the pH to 5 (near neutrality), the catalysts are still active but v_0 is lowered. Interestingly, the decrease of v_0 is smaller in the presence of light (50 and 75% for MS and NP respectively), than in the dark (59 and 85% for MS and NP respectively), and the MS catalyst exhibits a smaller loss of activity than the NP catalyst. For all the tests at pH=8, v_0 is reduced from more than 94%, indicating an important loss of activity. However, in all the range of pH, the best results are obtained under visible light. It is also important to note that the gains due to visible light seems to increase with the pH, v_0 being increased by factors of 1.5-1.6, 2-2.5, and 2.5-3, respectively at pH=3, 5 and 8. The existence of a relatively good catalytic activity at pH=5, may allow these catalysts to be used at neutral pH. These results suggest that γ -Fe₂O₃ NP and γ -Fe₂O₃/SiO₂ MS used as heterogeneous photo-Fenton catalyst have the potential to overcome the drawback of a narrow pH range of the conventional Fenton reaction.

3.3.2 Effect of the H₂O₂ dosage

The dependence of the decolorization efficiency of MO on the initial concentration of H₂O₂ ([H₂O₂]₀), was tested by changing [H₂O₂]₀ from 6.8×10⁻³ mol L⁻¹ to 0.68 mol L⁻¹.The

relationship between v_0 and $[\text{H}_2\text{O}_2]_0$ is shown in fig. 4B. In all cases, v_0 increases with $[\text{H}_2\text{O}_2]_0$, and the factors of increase are always of the same order (v_0 is multiplied by 1.7 to 2.2, from $[\text{H}_2\text{O}_2]_0 = 6.8 \times 10^{-3} \text{ mol L}^{-1}$ to $6.8 \times 10^{-2} \text{ mol L}^{-1}$, and by 2.7 to 4, from $[\text{H}_2\text{O}_2]_0 = 6.8 \times 10^{-2} \text{ mol L}^{-1}$ to 0.68 mol L^{-1}). This trend may be explained by the fact that the increase of $[\text{H}_2\text{O}_2]_0$ leads to an increase in the amount of HO^\bullet radicals, resulting in enhanced efficiencies for the removal of MO. The inhibition steps, such as reactions (10) and (11), which have been observed at high concentration of H_2O_2 , [16, 18] are not predominant in our case, although they may probably exist at a small extent.



Furthermore, the results prove that for both catalysts v_0 is always improved with visible light, independently of the initial amount of H_2O_2 , with slightly larger gains due to visible light for the low values of $[\text{H}_2\text{O}_2]_0$ (v_0 is increased by factor of 2.2-2.3, and 1.5-1.6, respectively at $[\text{H}_2\text{O}_2]_0 = 6.8 \times 10^{-3}$ and 0.68 mol L^{-1}). The lowest efficiency of visible light at high concentration of H_2O_2 may be correlated to a greater extent of the inhibitory steps (such as reactions (10) and (11)) under illumination. The most important point to retain is that both catalysts are still active at low H_2O_2 concentrations, especially under visible light, which has a great potential for an application in water-treatment, where a moderate amount of H_2O_2 is required for economical and technical reasons.

3.3.3 Effect of the amount of catalyst

The catalyst loading is one of the main parameters that can substantially affect the Fenton and photo-Fenton processes. Usually, it is expected that higher catalyst loadings lead to more rapid degradation of organic compounds. This can be explained by the fact that the rate of H_2O_2 decomposition increases with the amount of catalyst, which leads to a larger concentration of HO^\bullet radicals. In this study, the effect of the amount of catalysts was investigated by varying the equivalent concentration of iron from 0 to $3.65 \times 10^{-2} \text{ mol L}^{-1}$. The results for Fenton and photo-Fenton processes are shown in fig. 4C. It can be seen that in all the situations, v_0 linearly increases with the amount of iron. Whatever the amount of catalyst, better activities are obtained under irradiation. At high amount of catalyst, the ratio between v_0 under visible light and v_0 in the dark is quasi-independent of the amount of catalyst (the ratios vary between 1.5 and 2 for both catalysts). However, for the low amounts of catalyst,

larger ratios are obtained, especially with the MS catalyst. The better efficiency of visible light at low amount of catalyst can be analyzed in term of the relative importance of the uncatalyzed reaction under illumination. Indeed, assuming an order 1 with respect to the equivalent iron concentration, v_0 can be written as:

$$v_0 = k_{\text{obs}}[\text{Fe}]_{\text{T}} + v_{0,\text{uncat}} \quad (\text{Eq.4})$$

Where k_{obs} is the apparent constant of the catalyzed reaction (the slopes in fig.4C), $[\text{Fe}]_{\text{T}}$ the equivalent iron concentration, and $v_{0,\text{uncat}}$ the uncatalyzed initial rate (the ordinates in fig.4C).

The uncatalyzed reaction under visible light is not completely negligible, v_0 under visible light being twenty times larger than v_0 in the dark (see fig.2A and table S1). For the low values of $[\text{Fe}]_{\text{T}}$, v_0 is relatively small (especially for MS, the less active catalyst), and therefore, when visible light is applied, the contribution of $v_{0,\text{uncat}}$ is important, which mathematically increases the value of v_0 under light, in comparison to v_0 in the dark. Nevertheless, another explanation can be given for the larger efficiency of visible light at low amount of catalyst: the lower turbidity of the solution may enhance the transmission of visible light. However, we would expect to find in that case an absence of linearity between v_0 and the amount of catalyst. Finally, the most interesting feature is that the catalysts, when illuminated by visible light, maintain a good activity at very low amount of iron: despite the relative weakness of v_0 , complete decolorizations can be obtained in few hours for both catalysts, down to a value of $[\text{Fe}]_{\text{T}} = 5 \times 10^{-3} \text{ molL}^{-1}$, which corresponds to a $\text{H}_2\text{O}_2/\text{Fe}$ molar ratio of only 200/1.

3.3.4 Effect of the light intensity

We tested different commercial halogen lamps differing only in their nominal power which varied from 20W to 77W, on the photo-Fenton degradation of MO in presence of the NP catalyst. These halogen lamps essentially emits in the visible part of the spectrum (The temperature of the filament being at 2700K for all the lamps, similar emission spectra can be obtained), and it is well known that the visible light intensity linearly increases with its nominal power.

The experimental results presented in figure S6 clearly show that the rate of MO decomposition increases with the luminous intensity of the halogen visible lamp. Therefore when the intensity of the light is increased, both the initial rate v_0 , and the decolorization at 20

min, are improved. Although the increasing in efficiency seems to be slightly diminished for the larger nominal powers, best results are obtained with the 77W lamp, which corresponds to the stronger visible light intensity used in this study (a nominal power of 77W is equivalent to a visible light intensity of 1320 lumen according to the manufacturer). This demonstrates the possibility to increase the efficiency of the catalysts by increasing the light intensity of the lamp (the lamp used in our work being usually of 46W, which is equivalent to a visible light intensity of 700 lumen according to the manufacturer). The increase of the efficiency of the NP with the visible light intensity can be easily explained. As light intensity is increased, a larger number of photons are generated. More electron-hole pairs are then created at the surface of the γ -Fe₂O₃ NP. As a result, more HO[•] radicals are generated. This larger concentration in HO[•] radicals leads to a higher decomposition rate of MO.

3.4 Stability and reusability of the catalysts

Stability is an important property for an effective heterogeneous catalyst. One of the main limiting factors for the replacement of homogeneous Fenton catalysts by heterogeneous Fenton catalysts is the dissolution of a small part of iron ion during the depollution process, which may hinder the catalytic activity after several uses. The percentage of dissolved iron ions after 4H of reaction (Fe_L) was measured for both catalysts, in the absence and in the presence of light, for the three compounds (see table S1, columns 9 and 10 in the ESI part). The results show that leached iron detected in the reaction solution is minimal, with Fe_L less than 0.4%, (except in the case of the photodegradation of MB by the MS catalyst, where Fe_L is 0.7%), which corresponds to iron concentrations in the supernatant of less than 1.3×10^{-4} mol L⁻¹. Remarkably, no major increase of the leaching is noted when the catalysts are submitted to light, while this phenomenon seems to be independent from the catalyst. The slightly larger Fe_L observed for the MB pollutant with the MS catalyst may be due to the strong adsorption of this dye on this catalyst, which may remove additional iron from the surface of the γ -Fe₂O₃ NP, once the products resulting from the degradation of MB are desorbed.[35] These results, combined with additional experiments showing the low catalytic activity of the supernatant after 4h of reaction,[35] indicate that the contribution of the leached iron to the overall catalytic activity is weak. Moreover, the relatively low values of Fe_L is indicative of the good chemical stability of the γ -Fe₂O₃ NP during the reaction, which is required for the maintenance of their catalytic activity during several cycles. The stability of this catalyst is

also confirmed by the TEM and SAED-TEM images of the free NP after one photo-Fenton test on the model pollutant MO, and their dispersion in a HNO_3 10^{-3} mol L^{-1} aqueous solution (see figure S7 in the ESI). In comparison to the TEM image of the NP before catalysis (see figure 1-A), the TEM images of figure S7-A and B shows that the NP are characterized by similar sizes and morphologies. The TEM-SAED image of the sample (see figure S7-C) confirms that the nanoparticles are composed of $\gamma\text{-Fe}_2\text{O}_3$. The calibration of the SAED image allowed us to attribute the diffraction rings and spots to the interplanar spacings to the d_{440} , d_{400} , and d_{311} distances of the $\gamma\text{-Fe}_2\text{O}_3$ crystal lattice structure. Thus, the morphology and crystalline structure of the NP were not altered by the photocatalytic tests.

Reusability of a catalyst is of prime importance in assessing the practical application of a catalyst in water treatment. In practice, the reusability was only studied on the $\gamma\text{-Fe}_2\text{O}_3/\text{SiO}_2$ MS since this catalyst can be perfectly removed from the effluent after few minutes of settling over a magnet (see fig. 1C). On the contrary, our attempts to remove the $\gamma\text{-Fe}_2\text{O}_3$ NP by this method were unsuccessful (see fig. 1D), because of their high colloidal stability in water which also made difficult the other separation methods. The efficiency of the MS catalyst during five consecutive experiments was then investigated by regarding the decolorization of MO solutions. The catalytic tests were carried out for 4h, in the absence or in the presence of light. The results are shown in figure 5.

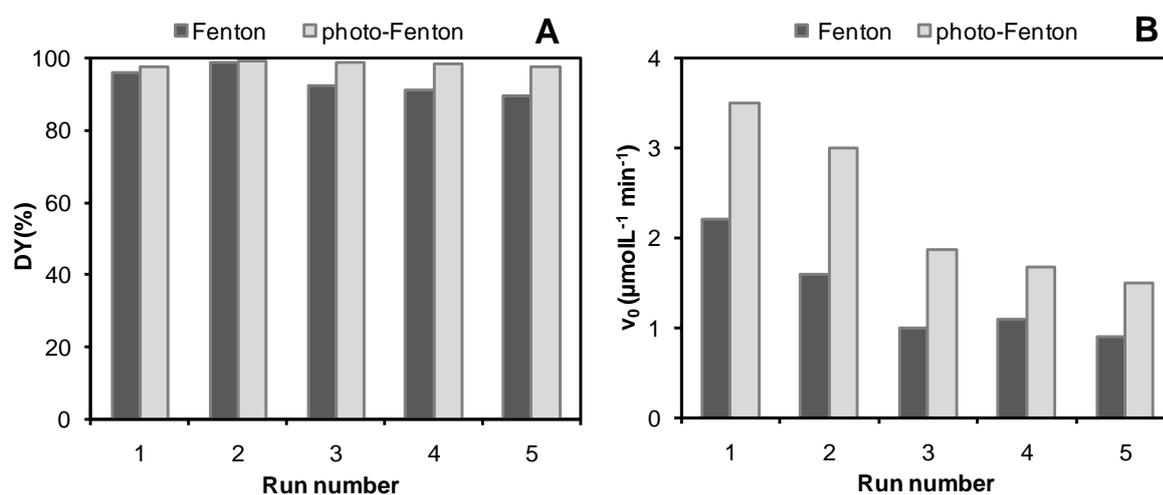


Fig. 5: Comparison of the reusability for five consecutive tests of the MS catalyst in the dark and under visible light, using MO as model pollutant. (A) Decolorization yields (DY) (taken at $t=4\text{h}$ and $t=3\text{h}$, for the Fenton, and the photo-Fenton processes, respectively). (B) Initial decolorization rates (v_0).

As can be seen in figure 5, the MS catalyst maintained a significant catalytic activity for at least five uses. For the photo-Fenton process, the MS catalyst retained a good activity after five repeated experiments, with complete MO conversions (97.5-99%), achieved within 3h (see fig. 5A), although v_0 was reduced from 57% (as shown in fig. 5B, v_0 vary from $3.5 \mu\text{mol L}^{-1}\text{min}^{-1}$ for the 1st cycle, to $1.5 \mu\text{mol L}^{-1}\text{min}^{-1}$ for the 5th cycle). Concerning the dark Fenton process, the catalyst also retained a relatively good activity, despite a slight decrease of the DY at 4h from 98% to 90%, and a reducing of v_0 from $2.2 \mu\text{mol L}^{-1}\text{min}^{-1}$ for the first cycle to $0.9 \mu\text{mol L}^{-1}\text{min}^{-1}$ for the 5th cycle, representing 59% of decrease. For each cycle, thanks to the visible light irradiation, v_0 is enhanced from a value oscillating between 1.5 and 1.9, indicating that the improvement factor due to light is almost constant. The moderate loss of activity as the catalyst is reused seems to be independent from the presence of the halogen lamp. This drop is not mainly due to the Fe leaching, taking into account the low amounts measured. Instead, we propose that the MS catalyst is progressively contaminated by some organic intermediates remaining strongly adsorbed on the iron oxide surface. For the tests in the dark, this has been qualitatively supported by the characterization of the catalyst after five cycles.[35] We have shown by SEM, N₂ volumetry, X-ray diffraction and UV-vis-NIR spectroscopy that the internal structure of the microspheres remain unchanged, despite a slight modification of their external morphology. Nevertheless, characterization of the microspheres by IRTF spectroscopy and thermogravimetric analysis has evidenced the presence of small amounts of organics.

4. Conclusion

The present work aimed to study the effect of a visible light emitting halogen lamp on the Fenton degradation of model organic pollutants, in presence of a heterogeneous catalyst. The catalyst was constituted by dispersed maghemite ($\gamma\text{-Fe}_2\text{O}_3$) nanoparticles (NP catalyst), or by the same nanoparticles encapsulated into microporous silica microspheres (MS catalyst). Both catalysts were characterized by a high magnetic susceptibility, and had therefore the potential to be easily recovered by a simple gradient of magnetic field. After their complete characterization, the catalysts were tested in the dark (Fenton Process), or under visible light illumination (photo-Fenton process), on three model organic pollutants, methyloange (MO), methylene blue (MB), and paranitrophenol (PNP). Much better catalytic activities were recorded in presence of visible light, whatever the catalyst, the pollutant tested, or the

experimental conditions. The large range of conditions studied in this work enabled us to suggest a mechanism for the visible light photodegradation. Since the level of v_0 increase due to visible light is poorly influenced by the nature of the pollutant, we proposed that the first steps of the degradation are due to photogenerated electron-hole pairs in the band gap of the $\gamma\text{-Fe}_2\text{O}_3$ nanoparticles. Illumination led also to a strong improvement of the mineralization of the pollutant, which is essential for the application of this process in water treatment. In addition, good activities were obtained in conditions which are considered to be unfavorable for the Fenton-type processes. The good decolorization rates obtained at neutral pH, at small amount of catalyst, or at small concentrations of H_2O_2 , through the use of a simple halogen lamp, demonstrated the possibility to extent the area of efficiency of the heterogeneous Fenton-type processes, and to reduce the associated operating costs. Furthermore, the study on the effect of the light intensity on the catalytic activity of the NP proves that it may be possible to improve again the efficiency of the catalysts by using a lamp with a higher nominal power. Another positive aspect is the experiments of long-term stability which showed that MS catalyst retained almost all of its catalytic activity after five repeated experiments under visible light. The good stability of this catalyst was also confirmed by the low level of iron leaching. In conclusion, we believe that these good results, due to the simple use of a low-cost, widely available and easy-to-handle halogen lamp will make possible the use of magnetic heterogenous Fenton catalyst in industrial wastewater treatment, especially for niche applications where recovery of the catalyst is a critical point.

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