Abatement of an Azo Dye on Structured C-Nafion/Fe-Ion Surfaces by Photo-Fenton Reactions Leading to Carboxylate Intermediates with a Remarkable Biodegradability Increase of the Treated Solution

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A novel C-Nafion/Fe-ion structured fabric capable of mediating Orange II decomposition in Fenton-immobilized photoassisted reactions is presented. The catalyst preparation requires the right balance between the amount of the Nafion necessary to protect the C-surface and the minimum encapsulation of the Fe-cluster catalytic sites inside the Nafion to allow the photocatalysis to proceed. The C-Nafion/Fe fabric can be used up to pH 10 under light to photocatalyze the disappearance of Orange II in the presence of H₂O₂. The photocatalysis mediated by the C-Nafion/Fe-ion fabric increased with the applied light intensity and reaction temperature in the reaction needing an activation energy of 9.8 kcal/mol. This indicates that ion− and radical−molecule reactions take place during Orange II disappearance. The build up and decomposition of intermediate iron complexes under light involves the recycling of Fe²⁺ and was detected by infrared spectroscopy (FTIR). This observation, along with other experimental results, allows us to suggest a surface mechanism for the dye degradation on the C-Nafion/Fe-ion fabrics. The C-Nafion/Fe-ion fabric in the presence of H₂O₂ under solar simulated light transforms the totally nonbiodegradable Orange II into a biocompatible material with a very high BOD₅/COD value. X-ray photoelectron spectroscopy (XPS) and sputtering by Ar⁺-ions of the upper surface layer of the C-Nafion/Fe-ion fabric allow us to describe the intervention of the photocatalyst down to the molecular level. Most of the Fe clusters examined by transmission electron microscopy (TEM) showed particle sizes close to 4 nm due to their encapsulation into the Gierke cages of the Nafion thin film observed by scanning electron microscopy (SEM) and optical microscopy (OM).

Introduction

The photo-Fenton reagent (Fe³⁺/Fe²⁺/H₂O₂) is an efficient and cheap reagent to generate oxidative radicals such as OH⁺ and HO₂⁻ in homogeneous reactions,† capable of oxidizing organic compounds to CO₂ and water. Iron in its ferrous and ferric form acts as a photocatalyst and requires a working pH ≤ 4. At higher pH values, iron precipitates as a hydroxide. The reversible iron cycle during a photoassisted Fenton homogeneous reaction involves the first step²

\[ \text{H}_2\text{O}_2 + \text{Fe}^{2+} \rightarrow \text{Fe}^{3+} + \text{HO}^- + \text{HO}^+ \quad k_1 = 40 - 60 \text{ M}^{-1} \text{s}^{-1} \quad (1) \]

In the second step, Fe³⁺-ions are reduced by light² (λ < 330 nm), which generates additional OPP radicals

\[ \text{Fe(OH)}^{2+} \xrightarrow{hv} \text{Fe}^{2+} + \text{HO}^+ \quad k_2 = 2 \times 10^{-3} \text{ M}^{-1} \text{s}^{-1} \quad (2) \]

The ligand to metal charge transfer (LMCT) via a complex with carboxylate radicals³ takes place in a redox photo-Kolbe process leading to decarboxylation of the organic compound

\[ [\text{RCOO} - \text{Fe}]^{2+} \rightarrow [\text{R}'] + \text{CO}_2 + \text{Fe}^{2+} \quad (3) \]

During the homogeneous Fenton reactions, the pH has to be adjusted twice, first to an acidic pH < 4 to carry out the Fenton pretreatment and then back to a neutral pH. In this way, it is possible to proceed with the lower cost second stage biological destruction of the organic compound(s) once the biocompatibility has been attained during the Fenton pretreatment. This is why supported Fenton-immobilized catalysts have been recently developed to overcome the costly pH adjustment on Nafion membranes,² silica structured surfaces,³ Nafion silica surfaces,⁴ polyethylene copolymers,⁵ and alginate microcapsules.⁶ These supported catalysts have been developed during the past few years with the intention of (a) withstanding the corrosion due to the oxidative radicals produced in solution, (b) fixing the Fe-ions on a support in a stable way so as not to allow them to leach out during the reaction, and (c) allowing the Fenton-mediated catalysis to proceed at acceptable kinetic rates.

Recently oven fibrous materials have become important as structured catalytic beds for reactor processes.⁷ C-fibers and

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fabrics have been used recently for metal deposition and tested as catalysts in a variety of processes. These materials are bidimensional and possess adequate mechanical and thermal resistance. The functional groups on C-fabrics can exchange extensively and retain strongly the Fe species necessary for the Fenton catalysis involving decomposition of H$_2$O$_2$.

In the present study, we present a C-Nafion/Fe-ion fabric as a composite material being applied for the first time as a stable immobilized catalyst decomposing Orange II by Fenton photo-assisted reactions. A novel composite catalyst consisting of C-filaments of 7 µm aligned in a woven structure covered by a Nafion film on which Fe-ions have been exchanged is studied in detail. The C-fabric presented in this study shows adequate characteristics during the oxidative Fenton processes when used in photoreactors with suitable dimensions and geometry for the objective of this study. The results presented show the photocatalytic decomposition of the model azo dye compound, the activation energy ($E_a$) of the process, the modeling of the reaction kinetics taking place in the cylindrical reactor, and the remarkable increase in the biodegradability of the pretreated solutions.

The C-structured loaded and unloaded surface characterization was carried out by electron microscopy (optical, SEM, and TEM), surface area measurements (BET), electrical resistance of the surface, photoelectron spectroscopy (XPS), and external reflection Fourier-transform infrared spectroscopy (ERFTIR) to determine the catalyst morphology and the surface composition at the molecular levels.

**Experimental Section**

**Chemicals and Fabrics.** Nafion perfluorinated ion-exchange resin 5% by weight solution in methanol (Aldrich 51, 921-1) was used as received. Acids and bases, methanol, Fe$_2$(SO$_4$)$_3$, 7H$_2$O, KSCN, Orange II, and H$_2$O$_2$ were p.a. Fluka AG products (Buchs, Switzerland) and used as received.

The C-fabrics were received from Longfellow Cambridge Ltd, Cambridge, U.K. After preliminary screening, a C-fabric 400 Tex with 6000 filaments/fiber of 7 µm each was chosen and used throughout this study on the basis of the favorable disappearance rate of Orange II. This C-fabric 400 Tex was 0.4 mm thick and 406 g/m$^2$ in weight and had a negligible surface area of 0.33 m$^2$/g. Other C-fabrics screened were 66 Tex and 200 Tex having 1000 and 3000 C-filaments/fiber. The reasons for the better performance of the C-fabric 400 Tex with regards to the 66 and 200 Tex C-fabrics will be dealt with in the Results and Discussion section. The C-fabrics used were all stable in air and room temperature. The electrical resistance found for the C-fabrics 66, 200, and 400 Tex were 8, 25, and 50 Ω/cm, respectively.

**Preparation of C-Nafion/Fe-Ion Fabrics.** The C-fabrics were cleaned with adequate amounts of methanol, immersed in the 5% Nafion solution, dried, washed with acid and H$_2$O$_2$, and subsequently exchanged with Fe$_2$(SO$_4$)$_3$·7H$_2$O. Finally, the C-Nafion/Fe-ion fabrics were washed with an abundant amount of distilled water and dried.

During the preparation of the membranes, the question of how many layers of Nafion were necessary to attain the most suitable photocatalytic disappearance of Orange II by the C-Nafion/Fe-ion fabric was investigated in a systematic way. It was found that one Nafion protective layer on the C-fabric was not enough. When two layers of Nafion were applied on the C-fabric, the performance of the C-Nafion fabric after Fe-exchange was still not reproducible. But when three layers of Nafion were applied, the results with the C-Nafion/Fe-ion fabric showed optimal Orange II decoloration kinetics. When four Nafion layers were applied prior to the Fe-exchange, the photocatalytically mediated disappearance of Orange II required much longer times. These less favorable decoloration kinetics were probably due to a thicker Nafion coating encapsulating the Fe-ions. With three Nafion coatings on the C-fabric the reproducibility found for the decoloration of Orange II was on the order of ±5%. This shows the validity of the preparative method used for the C-Nafion/Fe-ion fabrics used throughout this study.

**Photoreactor and Irradiation Procedures.** The photocatalytic activity of the structured C-Nafion/Fe-ion fabric was evaluated in 70 mL cylindrical reaction vessels. The light source was a Suntest solar simulator. The Suntest lamp had a wavelength distribution with about 7% of the photons between 290 and 400 nm. The emission profile of the photons between 400 and 800 nm followed the solar spectrum. The radiant flux of the Suntest light was measured with a power meter from LSI Corp., Yellow Springs, CO. The C-Nafion/Fe-ion fabric was located right behind the reactor wall. The H$_2$O$_2$ was added at the beginning of each run in the required amounts.

**Chemical Oxygen Demand (COD).** The measurement was carried out on a Hach DR/2000 optical unit. Test solutions (2 mL) were pipetted into a dichromate solution and digested at 150°C for 2 h. The optical density for the change in color of the dichromate solution was determined at λ = 430 nm.

**Biological Oxygen Demand (BOD).** This test was carried out by means of a BOD controller model 1020 (WTW). The unit was thermostated at 20°C. The pH of the samples was adjusted between 6.8 and 7.5 before an addition (10% v/v) of decanted sludge (inoculum from the biological treating station of Vidy, Lausanne) in the presence of the nutrients necessary for the bacterial activity.

**Analysis of the Irradiated Solutions.** Spectrophotometric analyses of the solutions were carried out using of a Hewlett-Packard 8452 diode-array spectrophotometer. The total organic carbon (TOC) was monitored via a Shimadzu 500 instrument equipped with an ASI automatic sample injector. The peroxide concentrations were assessed using Merkoquant paper and found to be at levels between 0.5 and 25 mg/L of H$_2$O$_2$. The dye disappearance was monitored by a Varian HPLC provided with a 9065 diode. A Phenomenex C-18 inverse phase column was used, and the dye signals were recorded at 229 and 254 nm. The gradient solution used was regulated with a buffer consisting of ammonium acetate and methanol.

**Fourier-Transform Infrared Spectroscopy.** The detailed description of the applied external reflection FTIR technique and procedure used has been previously reported. The infrared reflection spectra of catalyst samples were recorded on a Bruker IFS55 FTIR spectrometer equipped with an MCT detector and a reflection attachment (Seagull). A wire-grid polarizer was placed before the sample and provided p- or s-polarized light. These accessories were from Harrick Scientific Co. It was found for the system under investigation that the most informative reflection spectra could be recorded by the use of p-polarized light and an angle of incidence of 20°. The unit of intensity was defined as $-\log(R/R_0)$, where $R_0$ and $R$ are the reflectivities of the catalyst sample after the reaction and before the reaction directly after its preparation, respectively. Both sample and reference spectra are averaged over the same number of 1000 scans.

**X-ray Photoelectron Spectroscopy (XPS).** Measurements were carried out in a Leybold-Heraeus instrument calibrating the binding energies (BE) to the Au f$_{7/2}$ level taken as 84.0 eV. The evaluation of the binding energies of the embedded Fe
clusters was carried out following the DIN norms\textsuperscript{10} and, after the sensitivity factors were taken into consideration, allowed a reproducibility of $\pm 5\%$ in the measurements. The quantitative evaluation of the experimental data was carried out with a Shirley-type background correction\textsuperscript{11} due to the electrostatic charging of the particles during the measurements. The sputtering of the external surface loaded fabric was carried out with Ar-ions, and after the Ar-ion bombardment, the recorded oxidation states of the surface species may not entirely correspond to the true states in the C-Nafion/Fe-ion fabric because the Ar-ion treatment changes the nature of the fabric to some extent.

### Optical Microscopy, Transmission Electron Microscopy (TEM), and Scanning Electron Microscopy (SEM)

The optical microscope, an Olympus 9M, provided with a digital camera was used to record the images of the C-fabrics and the largest Nafion deposits at enlargements of 24, 40, and 66 times. The scanning electron microscopy (SEM) was performed with a JEOL 35 CF (15 kV, 15 nm point resolution) to observe the C-fabrics covered by some Nafion deposits with a gold layer 0.1 nm thick. A field emission TEM, the Philips CM 120 microscope, was used to observe the Fe clusters on the Nafion on the C-fabric surface. The C-fabric was coated with epoxy resin, and the fabric was cut with a microtome to a thin layer of 50 nm at a 90° angle for the experimental observations. The instrument used for electron microscopy (120 kV, 0.35 nm point resolution) was equipped with an energy-dispersive X-ray analysis EDAX to identify the Fe clusters on the support.

### Results and Discussions

#### Immobilized C-Nafion/Fe-Ion Fabrics Versus Fenton Homogeneous Photoassisted Decoloration of Orange II

Figure 1 (trace a) shows the lack of decoloration of Orange II in the dark in the presence of a C-Nafion/Fe-ion fabric. Under Suntest light irradiation, Figure 1 (trace b) shows that Orange II is completely decomposed in $\sim 40$ min. After a photocatalytic run the amount of Fe$^{3+}$ was analyzed in solution by complexing the Fe$^{3+}$ with thiocyanate and found to be always $\approx 0.5$ mg/L. To reference the results obtained (trace c), Orange II was run under Suntest irradiation with a solution containing Fe$^{3+}$ (0.5 mg/L). The results in Figure 1 show that the decoloration of Orange II within 40 min is mainly due to the C-Nafion/Fe-ion fabric and only to a very small extent to the Fenton homogeneous residual reaction going on in parallel in the reactor.

Figure 2 shows the practical absence of Orange II decoloration when C-fabrics, C-Nafion fabrics, and C-Nafion/Fe-ion fabrics were used as catalysts in the dark or light-induced processes in the absence of H$_2$O$_2$. A small decoloration was observed during Orange II decoloration in the dark due to some minor adsorption or interaction between the Orange II and the redox impurities of the C-fabric surface or both. Orange II adsorption was not observed on C-Nafion and C-Nafion/Fe-ion fabrics.

Up to eight recycles were carried out to test the catalytic nature of the Orange II mediated decoloration by C-Nafion/Fe-ion. The experimental conditions used were the same as those in Figure 1. After each recycling, the fabric was thoroughly washed with distilled water.

#### Kinetics of Orange II Decomposition as a Function of the Initial pH and H$_2$O$_2$ Concentration

Figure 3 shows Orange II disappearance during Suntest irradiation mediated by C-Nafion/Fe-ion fabrics as a function of the initial pH of the solution. Slower kinetics for processes at initial pH values of 10, 8.1, 7.0, and 5.1 compared to pH 2.7 is observed. Similar results have been observed before for Orange II degradation in homogeneous photoassisted Fenton reactions. The complex
shape for the Orange II concentration as a function of time suggests the presence of pH-dependent transient intermediates affecting the course of Orange II disappearance. These solutions did not precipitate, and Fe aqueous complexes intervening favorably during Orange II disappearance begin to be effective after ~15 min for Orange II solutions with an initial pH > 5.1. The insert in Figure 3 shows the decrease of TOC for Orange II solutions with initial pH of 2.7. TOC reduction of Orange II solutions was observed only at this low initial pH-value. This implies that intermediates required for a reduction of TOC are only effective at acidic pH values ≤ 4.2,12,13 If we start at higher pH values, the intermediates produced in solution preclude further degradation of Orange II in solutions with an initial pH ≥ 5.1 as seen in Figure 3. The residual TOC noted in the insert of Figure 3 did not decrease with increasing concentration of H₂O₂. Because decoloration proceeds well at pH values up to 10 (Figure 3), the intermediates leading to decoloration are quite different in nature than the ones involved in the TOC reduction during the photocatalysis of Orange II as reflected in the rates

Orange II $\overset{k_d}{\rightarrow}$ organic intermediates $\overset{k_{oa}}{\rightarrow}$ CO₂

where $k_d \gg k_{oa}$ (4)

The OH$^+$ and HO₂$^*$ radicals react rapidly and unselectively with aromatic compounds with rate constants of $10^9 - 10^{10}$ M⁻¹ s⁻¹ and $10^6 - 10^8$ M⁻¹ s⁻¹, respectively. Intermediates produced during Orange II degradation under light irradiation in the presence of O₂ and H₂O₂ have been reported before, but the exact nature of their composition in the case of homogeneous Fenton systems is not known. By HPLC, it was not possible to determine these Fe complex intermediates present in solution, and GC-MS work is under way to clear this point. The pH for Orange II solutions with an initial pH above 5.1 reaches a pH of ~4 after 15 min of light irradiation. This time period coincides with the start of the decoloration reported for Orange II solutions in Figure 3. Only when the pH value was close to 4 was the decoloration of Orange II able to proceed as reported in Figure 3. At more basic pH values, the Fe(H₂O)₆³⁺ species (pK₆ = 2.8) deprotonates rapidly to Fe(H₂O)₅(OH)²⁺ and Fe(H₂O)₄(OH)⁺ 12–15 hindering the decoloration of Orange II at reaction times <15 min (Figure 3). In addition at acidic pH values, the OH$^+$ radical has been reported to be the dominant species when Fe(OH)²⁺ is present in solution.16,17 Recently, adduct formation between OH$^+$ and Orange II has been reported during Orange II decomposition at acidic pH values involving protonated species.18,19

Figure 4 presents the decoloration kinetics of Orange II as a function of the added concentration of H₂O₂. It is readily seen that increasing the amount of H₂O₂ leads to a faster Orange II decoloration up to 10 mM. Beyond the latter H₂O₂ concentration, an increase in the amount of H₂O₂ added slows down the decoloration because the propagation step shown in eq 5 is hindered by the excess H₂O₂ scavenging the available OH$^+$ radicals

$$\text{H}_2\text{O}_2 + \text{OH}^+ \rightarrow \text{H}_2\text{O} + \text{HO}_2^* \quad k_5 = 1.2 \times 10^7 \text{M}^{-1} \text{s}^{-1}$$ (5)

Energy of Activation ($E_a$) and Light Intensity Dependence of the Orange II Decoloration. Figure 5 shows the decoloration of an Orange II (0.2 mM) solution as a function of the applied temperature. It is readily seen from Figure 5 that an increase in temperature has a beneficial effect on the decoloration rate. The $E_a$ was determined by following the concentration of Orange II at 20, 40, 55, and 70 °C vs time. At each temperature, the initial decrease of the decoloration rate in millimoles per minute was estimated. Only the initial values of the tangent in Figure 5 were taken into account for the calculation of $k$ because of the shapes observed for the decoloration process in Figure 5. In this way, the input due to the light energy is minimized and the temperature effect is considered in Figure 5. A value of 9.8 kcal/mol was obtained from the graph of ln $k$ vs 1/T K. The latter value suggests that ion–molecule reaction and radical–molecule reactions requiring some activation energy occur during Orange II disappearance, as well as the radical–radical reactions that need no $E_a$.

Figure 6 shows the dependence of the decoloration on the intensity of the applied light from the Suntest solar simulator. The intensity of 50 mW/cm² corresponds to 50% of the full light intensity of equatorial solar radiation perpendicular to the equator (AM1). Consequently, 90 mW/cm² corresponds to 90% of AM1. Accelerated decoloration was obtained when a medium-pressure mercury arc (Applied Photophysics, U.K.) with the main radiation centered at 366 nm was used as the radiation source with $6 \times 10^{10}$ photons/s. A quicker decoloration was obtained with 90 mW/cm² as compared to a light intensity of 50 mW/cm². This implies that there is no saturation of the Orange II solution by the photon density of the applied light. Orange II, being the light absorber under visible irradiation, initiates the decoloration process mediated at a later stage by C-Nafion/Fe-ion fabrics in this dye-sensitized process.
Decoloration of Orange II by a C-Nafion/Fe-ion fabric. Figure 7 presents the degradation kinetics for two concentrations of Orange II under Suntest light irradiation in a cylindrical reactor stirred at a rate that avoids mass transfer problems despite the vigorous stirring employed. The shape of the curves in Figure 7 suggest lateral boundaries in the reactor vessel due to the C-Nafion/Fe-ion fabric having a small active zone. In this zone, the oxidative radicals are generated. The chemical reaction between Orange II and the oxidative radicals occurs under a turbulent flow regime in the reaction vessel.

Increase in the Biodegradability of Orange II During the Pretreatment on C-Nafion/Fe Fabrics. Figure 8a shows the BOD$_5$ evolution of untreated and treated solutions of Orange II (0.2 mM) showing the beneficial effect of the photocatalysis mediated by C-Nafion/Fe-ion fabrics on the biodegradability of the solution. As shown in the lower part of Figure 8a, the initial Orange II solution is nonbiodegradable. But when Orange II is pretreated for 90 and 120 min, the intermediates present a very high biodegradability. On smaller organic intermediates with a higher degree of oxidation, the bacterial action proceeds more favorably, as reported for other model compounds from our laboratory. The photohydroxylation and photooxidation of the aromatic ring make the bacterial attack easier because the resonance energy of the ring has been lowered, and the bacteria benefits additionally from the energy released because of this bond breaking.

Figure 8b shows the evolution of the BOD$_5$/COD ratio as a function of pretreatment time. This ratio increases up to 0.35 and 0.76 after 90 and 120 min of pretreatment. As a reference, the parameter for biodegradable wastewaters is 0.4. The insert in Figure 8b shows the evolution of the average oxidation state (AOS) as a function of time. The AOS represents the oxidation state of a solution as $+4$ for CO$_2$ and $-4$ for CH$_4$. The insert in Figure 8b indicates that with reaction time the intermediates of Orange II decomposition become more oxidized during the pretreatment.

Fourier-Transform Infrared Spectroscopy of C-Nafion/Fe-ion Fabrics. Figure 9 shows the external reflection FTIR spectra of a C-Nafion/Fe-ion fabric after reaction with Orange II for different times. The spectrum of the “fresh” catalyst directly after preparation was used as reference. The observed positive bands at 1670 and 1640 cm$^{-1}$ indicate the presence of surface carboxylate groups. The amount of these surface groups is the highest after 30 min (Figure 9b) of the Orange II photodecomposition reaction, whereas at the beginning and after 50 min of reaction, the surface densities of the carboxylate groups are very low. The broad negative absorbance band at around 1620 cm$^{-1}$ is assigned to water molecules surrounding iron ions in the Nafion structure. This water is replaced by the adsorbed decomposition products with carboxylic and probably aromatic groups during photodegradation of Orange II.

Figure 10 shows the reflection infrared spectra of the C-Nafion/Fe-ion fabric equilibrated with Orange II in dark after different conditioning times. The observed band at around 1620 cm$^{-1}$ is clearly visible in Figure 9a,c, is assigned to water molecules surrounding iron ions in the Nafion structure. This water is replaced by the adsorbed decomposition products with carboxylic and probably aromatic groups during photodegradation of Orange II. Figure 10 shows the reflection infrared spectra of the C-Nafion/Fe-ion fabric equilibrated with Orange II in dark after different conditioning times. The observed band at around 1620 cm$^{-1}$ is clearly visible in Figure 9a,c, is assigned to water molecules surrounding iron ions in the Nafion structure. This water is replaced by the adsorbed decomposition products with carboxylic and probably aromatic groups during photodegradation of Orange II.
Figure 11a shows the external reflection infrared spectra of the C-Nafion/Fe-ion fabric after equilibration with different concentrations of oxalic acid. The spectrum of the C-Nafion/Fe-ion fabric after preparation is our reference. Oxalic acid at 1 mM concentration forms a significant amount of complexes with iron present in the Nafion. This is indicated by the presence of two strong absorbance bands: a shoulder at 1688 cm\(^{-1}\) and a band at 1646 cm\(^{-1}\). With a higher oxalic acid concentration in solution above 5 mM, iron is removed from Nafion. The broad negative absorption bands at around 1620 cm\(^{-1}\), clearly visible in traces b and c of Figure 11a, indicate removal of water molecules surrounding iron ions in the Nafion structure. This observation supports the conclusion about the strong leaching of iron from Nafion with a concentrated oxalate solution.

Figure 11b shows external reflection spectra of the C-Nafion/Fe-ion fabric after exposure to an oxalic acid solution for 16 h and then in reaction with Orange II. The spectra of the C-Nafion/Fe-ion fabric after exposure to different oxalic acid solutions for 16 h were used as the references to see the effect of Orange II decomposition. At 1 mM oxalate concentration and 2 h of irradiation of the Orange II solution (trace a, Figure 11b) the reflection spectrum shows two negative bands: a shoulder at 1690 and a strong band at 1646 cm\(^{-1}\). The band positions and intensities are almost identical to those presented in Figure 11a, but they are negative. This indicates the disappearance of the produced iron oxalate complex during exposure to 1 mM oxalate solution after the photodegradation of Orange II. The C-Nafion/Fe-ion fabric sample exposed to 5 and 10 mM oxalate solution removed the majority of the iron from the Nafion, and therefore, oxalic acid was adsorbed in an almost negligible amount (traces b and c, Figure 11b). As a consequence, after Orange II photodecomposition for 2 h, the catalyst shows only a very small removal of the carboxylate surface species.

Figure 11c shows the results for the photodegradation of Orange II under light in the presence of C-Nafion/Fe-ion fabrics...
Figure 12. Fe species as a function of the erosion depth (a) during the Ar-ion sputtering of the C-Nafion/Fe-ion fabric surface, (b) concentration gradient of the different species of fluoro compounds as a function the erosion depth during Ar-ion sputtering of the C-Nafion/Fe-ion fabric surface, and (c) Removal of the surface from C–F groups of the C-Nafion/Fe-ion fabric surface with increasing erosion of the catalyst topmost layers until reaching the stable profile of the C concentration at a certain depth inside the C-Nafion fabric.
that have been pretreated in two different ways. Trace a represents the photodegradation of Orange II solution on the C-Nafion/Fe-ion fabric that was pretreated in 10 mM oxalic acid for 2 h and then was washed and used as a photocatalyst. Trace b shows the photodegradation of the same Orange II solution but with a photocatalyst pretreated for 2 hours in a 1 mM oxalic acid solution. The latter pretreatment of the photocatalyst did not dissolve the Fe-ions from the Gierke cages of Nafion, and the Orange II decomposition preceded readily. The results presented in Figure 11c are entirely consistent with the results obtained by FTIR (Figure 11a,b) supporting the conclusion about significant reactivity of the C-Nafion/Fe-ion fabrics in solution under 5 mM acetic acid.

Another important finding of the IR spectroscopy is that the main decomposition channel of Orange II on a C-Nafion/Fe-ion fabric involves an Fe-carboxylate redox photo-Kolbe process leading to the decarboxylation of the organic compound as mentioned previously in eq 3.

**X-ray Photoelectron Spectroscopy (XPS).** No significant difference was observed in the oxidation state of the Fe present in the C-Nafion/Fe-ion fabrics before and after photocatalytic decomposition of Orange II. A Gaussian–Lorentzian fitting of the XPS peaks was carried out by the Shirley-type background correction, subtracting the X-ray satellite peaks. Polynomial second-order fits of the experimental curves were directed toward the matching of the asymmetric shape of the corrected XPS signals referencing to the known binding energies: Fe metal 706.7 eV; FeO 709.6 eV; Fe2O3 710.9 eV; Fe3O4 711.4 eV. Sputtering of the C-Nafion/Fe-ion fabrics was carried out with Ar and the XPS depth profile and was recorded on 3D profile graphical plots that are shown below.

Figure 12a presents the continuous 3D representation of the Fe surface concentration. It is clearly seen that the maximum Fe concentration is below the topmost layers (of ~2 Å each) shown for the Fe2p3/2 doublet centered at 714.7 eV in Figure 12a. The estimation of 2 Å for each layer is an approximation.

![Figure 13. Scanning electron microscopy of (a) a C-woven fabric (400 Tex, each fiber is made up of 6000 filaments of 7 μm diameter each) and (b) one filament 7 μm in diameter making up the basic fibers used in the C-fabric (400 Tex).](image)

When erosion of the C-Nafion/Fe-ion fabric was carried out by sputtering with Ar+-ions, the layers of Nafion and of the C-fibers were stripped at a different rate. This leads to a different stripping depth for Nafion and C. The absence of Fe from the topmost layers is due to two factors: (a) the Fe-ions exchanging with the H+ of sulfonic pending group inside the Gierke cages placing the Fe-ions at some distance from the topmost Nafion surface and (b) the selective shielding by C-F groups of the Fe species in the interior of the Nafion Gierke cages.

Figure 12b presents the concentration gradient of F (for the different varieties of fluoro compounds found on the fabric surface) showing that the Nafion F surface species decrease in the layers located further away from the fabric surface. This shows that the Nafion forms a good film on the C-fabric. Furthermore, a second shoulder appears as the erosion penetration depth increases revealing a second and different C-F compound at some depth from the fabric surface. The removal of the C-F groups is seen to proceed with increased sputtering time leading to a zone where the C-substrate concentration predominates. The number of layers removed in terms of the C-F species in Figure 12b matches the erosion zone for the C-F species reported in Figure 12c. Figure 12e backs quantitatively the results presented in Figure 12b. The C-F species is the main component of the Nafion polymer, and it is the decomposition of H2O2 over the Fe-ionic species in the Nafion cages that produces the oxidative radicals necessary for Orange II decomposition. Furthermore, experiments following the sulfur enrichment of the C-Nafion/Fe-ion fabrics show that the topmost layers removed by Ar+-ion
sputtering at 165/165 eV are within the same distance from the surface as the C–F layers observed for the Nafion sulfonate with a binding energy value (BE) of 1697/170 eV.

Microscopy Studies on the C-Fabrics and the Loaded C-Fabric. A detailed presentation of the uneven surface of the C-fabric comprising the bundle of C-fibers is presented by

Figure 15. Optical microscopy (a) of the C-Nafion/Fe-ion fabric surface after Orange II photocatalytic decoloration and (b) transmission electron microscopy of the homogeneously dispersed Fe clusters on the C-Nafion/Fe-ion fabric surface before the photocatalytic runs on Orange II. A media size of 3.8 nm was accounted for from the histogram of the sizes of the Fe clusters on the C surface.
scanning electron microscopy (SEM) in Figure 13a and a single C-filament is shown in Figure 13b. The scale found in the lower right-hand in Figure 13b shows that each individual filament has a diameter of 7 μm. Orange II disappearance in the active zone near the C-Nafion fabric where the oxidative radicals are generated by H2O2 on the Fe clusters was optimized and proceeds in 40 min (Figure 1). The voids in the C-fabric structure would not be beneficial with regard to a regular and continuous generation of oxidative radicals on the C-fabric in the presence of H2O2.

During the dip coating deposition of Nafion onto the C-fabric, it reveals submicron and micron Nafion deposits as shown in Figure 14a,b by scanning electron microscopy. Submicron-sized thin Nafion deposits were observed on the 400 Tex C-fabric and on the C-filaments covering a large section surface area. This is shown in Figure 14a. Thicker Nafion deposits of 20–30 μm were also observed along with the former ones covering 10–20% of the C-fabric and C-filaments as shown in Figure 14b.

Figure 15a presents the C-Nafion/Fe-ion fabric with the detailed Nafion deposits protruding out of the surface of the C-filament. The Nafion deposits containing the Fe-ions stay on the C-filaments after several cycles even up to runs at pH 10. The surface of the C-filaments was still covered with thin Nafion submicron deposits revealing the improved stability of the C-Nafion/Fe-ion fabrics with respect to the glass-Nafion/Fe submicron deposits revealing the improved stability of the C-filaments after several cycles even up to runs at pH 10. A detailed Nafion deposits protruding out of the surface of the C-filament is shown in Figure 13b. The scale found in the lower right-hand in Figure 13b shows that each individual filament has a diameter of 7 μm. Orange II disappearance in the active zone near the C-Nafion fabric where the oxidative radicals are generated by H2O2 on the Fe clusters was optimized and proceeds in 40 min (Figure 1). The voids in the C-fabric structure would not be beneficial with regard to a regular and continuous generation of oxidative radicals on the C-fabric in the presence of H2O2.

This work presents the first report on C-Nafion/Fe-ion fabrics that as active supported photocatalysts in Fenton immobilized reactions. The optimization of the decomposition of Orange II taken as a model compound has been worked out in detail. The mechanism of the decoloration of azo dye was backed by experimental evidence obtained by infrared spectroscopy showing the formation and disappearance of the carboxylates intervening in the photocatalytic process. From the result obtained by FTIR spectroscopy, the decomposition channel of Orange II on the C-Nafion/Fe-ion fabric involves an Fe–carboxylate redox photo-Kolbe reaction. A detailed characterization of the surface parameters of the C-Nafion/Fe-ion fabric has been worked out through XPS, SEM, TEM, and BET techniques. A remarkable increase of the Orange II biodegradability was observed after the Fenton immobilized photoassisted pretreatment.

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References and Notes


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