

Degradation of the pharmaceuticals nimesulide and ibuprofen using photo-Fenton process: toxicity studies, kinetic modeling and use of artificial neural networks

Degradação dos fármacos nimesulida e ibuprofeno empregando processo foto-Fenton: estudos da toxicidade, modelagem cinética e emprego de redes neurais artificiais

Rubens Teles Monteiro, Rayany Magali da Rocha Santana, Ana Maria Ribeiro Bastos da Silva, Alex Leandro Andrade de Lucena, Léa Elias Mendes Carneiro Zaidan, Valdinete Lins da Silva, Daniella Carla Napoleão

Departamento de Engenharia Química, Universidade Federal de Pernambuco, PE, Brasil

rubens.engenheiroquimico@gmail.com; rayanymagalirocha@gmail.com; amrbsilva@ig.com.br; lucenaalex3@gmail.com; leazaidan@yahoo.com.br; leaq_val@yahoo.com.br; danicarlan@gmail.com

Abstract

*The growth of pollution in aquatic environments increases every day, causing compounds like pharmaceuticals to be detected in surface waters. Thus, techniques such as advanced oxidation processes (AOP) have been used to degrade these compounds. In this work, the efficiency of AOP in the degradation of nimesulide and ibuprofen pharmaceuticals was evaluated through chromatographic analysis as well as organic matter through the levels of chemical oxygen demand (COD) and total organic carbon (TOC). It was verified that the photo-Fenton process presented the best results, degrading 89.70% of nimesulide and 93.35% of ibuprofen. This same process managed to reduce COD by 91.60% and mineralize 90.04% of the TOC. The kinetic study showed a good linear fit ($R^2=0.993$) for the clustered kinetic model, as well as a good fit to the mathematical model of artificial neural networks (ANNs), with a value of $R^2=1.000$ for the MLP4-4-1 BFGS 4567 model. Finally, the toxicity of the solution after treatment was verified against the seeds of *Lactuca sativa*, *Cichorium endivia*, *Ocimum basilicum* and American Hard grain. It was found that the seeds that received the solution before treatment had a lower germination amount than the ones where the post AOP treatment solution was added. Then, the root growth was evaluated, in which a relative toxic effect was observed.*

Keywords: *Ibuprofen; Nimesulide; Photo-Fenton*

Resumo

*O crescimento da poluição de ambientes aquáticos tem aumentado todos os dias, fazendo com que compostos como os fármacos sejam verificados em águas superficiais. Desse modo, técnicas como processos oxidativos avançados (POA) tem sido utilizadas. Neste trabalho a eficiência dos POA na degradação dos fármacos nimesulida e ibuprofeno foi avaliada, através de análises cromatográficas, bem como de matéria orgânica através dos níveis de demanda química de oxigênio (DQO) e carbono orgânico total (COT). Verificou-se que o processo foto-Fenton apresentou os melhores resultados degradando 89,70% do nimesulida e 93,35% do ibuprofeno. Esse mesmo processo conseguiu reduzir em 91,60% a DQO e mineralizar 90,04% do COT. O estudo cinético mostrou bom ajuste linear ($R^2=0,993$) para o modelo cinético agrupado, além de uma boa adequação ao modelo matemático de redes neurais artificiais (RNA), com um valor de $R^2=1,000$ para o modelo MLP4-4-1 BFGS4567. Por fim, verificou-se a toxicidade da solução após tratamento, frente às sementes de *Lactuca Sativa*, *Cichorium endivia*, *Ocimum basilicum* e do grão Americano Hard. Verificou-se que as sementes que receberam a solução antes do tratamento apresentaram uma quantidade menor germinação, do que quando foi adicionada a solução pós-tratamento via POA. Em seguida, avaliou-se o crescimento radicular, no qual foi percebido relativo efeito tóxico.*

Palavras-chave: *Ibuprofeno; Nimesulida; Photo-Fenton*

1 INTRODUCTION

Different organic pollutants have been studied with regard to their persistence in the environment. In water the concern is even greater, since this resource is more and more scarce when it comes to drinking standards. Thus, different treatment processes have been studied, since conventional methods of wastewater treatment (physical-chemical and biological) are not able to promote the degradation of more persistent compounds such as pharmaceuticals (NAPOLEÃO et al, 2015) polycyclic aromatic hydrocarbons (ROCHA et al, 2014), plasticizers, pesticides, hormones and phenolic compounds (ZAIDAN et al, 2016).

In order to detect these pollutants in aquatic matrices, and at different levels of concentration, the effectiveness of treatments such as advanced oxidation processes (AOP) has been evaluated (SANTOS-JUANES et al, 2011), as well as the adsorption processes (RAJABI et al, 2016). Different types of AOP have shown good results, since they act by oxidizing the organic compounds, through the release of the hydroxyl radicals, which lead to the mineralization of the organic matter releasing inorganic compounds, CO₂ and H₂O. Among the most used AOP are the Fenton reaction, which occurs in the presence of Fe²⁺ ions, combined with a strong oxidizing agent, such as hydrogen peroxide (H₂O₂); The photo-Fenton process, which in addition to the Fenton reagents employs ultraviolet (UV) or visible radiation; the UV/H₂O₂ action, in which there is only the combination of H₂O₂ and radiation; and photolysis, that involves, only, the use of radiation (ARAÚJO et al, 2016). These processes act differently, depending on, among other factors, the pollutant in question.

In the case of the degradation of pharmaceuticals, studies have revealed that the processes involving the presence of radiation in conjunction with H₂O₂ present more satisfactory results (CARLSON et al, 2015; ALHARBI et al, 2017; NAPOLEÃO et al, 2018). This is because the radiation action leads to the photo-decomposition of this reagent, which generates hydroxyl radicals capable of promoting the oxidation of highly complex organic pollutants (NAVARRO; GABALDÓN; GÓMEZ-LOPEZ, 2017; LIMA; ALMEIDA; PAULA, 2016). Such decomposition mechanism is described in Equations 1 to 3.



The oxidizing power of hydroxyl radicals is often increased with the help of iron ions, which act to catalyze the reaction (BRITO; SILVA, 2012). In the photo-Fenton process, the hydroxyl radicals act to regenerate Fe³⁺ ions, which undergo a process of reduction (COSTA et al, 2014) according to Equation 4.



Studies on the combined performance of conventional treatments with AOP processes (Fenton process) to the degradation of phenol were able to promote a degradation of 99.7% of the compound, which presents in a recalcitrant way in aquatic matrices (SKORONSKI et al, 2015). In the same sense,

the photo-Fenton process has also been shown to be efficient in the removal of pharmaceutical contaminants present in water. Giri and Golder (2015) observed the application of this type of treatment to the drug dipyrone and obtained a degradation of 96.40%. Romero et al (2016) and Funai et al (2017) achieved 100.00% degradation by applying the same process to degrade the magnesium and sodium valproate drugs in aqueous medium.

In order to evaluate the percentage of degradation of the most diverse pollutants, it is necessary to have adequate analytical techniques. Among the most used ones is the chromatography, being the high-performance liquid chromatography (HPLC) used in the identification and quantification of pharmaceuticals (BIALK-BIELINSKA et al, 2016; PEAKE et al, 2016). To ensure greater reliability of the results, the methodology validation has been used, so that the analyzes can be said to be precise and accurate (ZANCHETA; PENA; GONÇALVES, 2015). Regarding the determination of organic matter levels, two parameters are the most applied, the chemical oxygen demand (COD) and the total organic carbon (TOC) (LUAN et al, 2017). It is based on the evaluation of these parameters that the kinetic monitoring of the degradation of pharmaceuticals by AOP is carried out. Based on that, the time in which the pollutants are mineralized can be determined (NAPOLEÃO et al, 2013).

Although advanced oxidative processes are well known and reported in the literature, it is sometimes difficult to clearly state the influence of the reaction parameters, especially when it is desired to apply such processes on an industrial scale. Based on this, some authors have made use of artificial neural networks (ANNs) to provide predictive models capable of adjusting nonlinear data of complex systems (JALIL et al, 2014; MORAES et al, 2016). Moraes et al (2016) state that a photo-oxidation process can be described efficiently through an ANN using the experimental data of total organic carbon as the output variable.

In addition to evaluating the parameters involved in AOP, their kinetic models and efficiency in the degradations of compounds considered to be persistent, recent studies show that just promoting the degradation is not enough. It is essential too, to carry out experiments evaluating the toxicity of the intermediates formed during the treatment (NAPOLEÃO et al, 2018). In this sense, researches have been carried out with the purpose of evaluating the environmental risks caused by the pollutants, as is the case of pharmaceuticals, compounds widely used by the population, which are often discarded in an improper way (VALCÁRCEL et al, 2011; PINTO et al, 2016).

Among the various classes of drugs studied are the anti-inflammatories, which includes nimesulide and ibuprofen. Nimesulide also has antipyretic and analgesic properties and is used in the treatment of febrile conditions, inflammatory processes related to the release of prostaglandins, besides being prescribed as an analgesic for dental infections and postoperative pain (SILVA; MENDONÇA; PARTATA, 2014). Ibuprofen is indicated for the relief of fever, trauma, headache, muscle and joint pain, as well as dental inflammation, sore throat and other symptoms associated with colds and flu (STOYANOVA, VINAROV, TCHOLAKOVA, 2016). These drugs are, therefore, widely used by the population, and somehow returned to the environment through the excretion process. In addition, they reach the environment coming from the production process itself, through the disposal of wastewater treatment plants of pharmaceutical industries (CHRISTOU et al, 2017; EBELE et al, 2017).

In this context, the present work aims to evaluate the efficiency of the degradation of nimesulide and ibuprofen in aqueous solution using different types of advanced oxidation processes. The study also sought to evaluate the most efficient AOP degradation kinetics, to use mathematical modeling using artificial neural networks and to verify the toxicity of the reaction intermediates against the

seeds of *Lactuca Sativa* (lettuce), *Cichorium endivia* (chicory), *Ocimum basilicum* (basil) and *Americano Hard* grain (wheat).

2 MATERIAL AND METHODS

2.1 Identification and quantification of the pharmaceuticals nimesulide and ibuprofen

An analysis of the drugs under study was carried out in high performance liquid chromatography (HPLC) equipment (Shimadzu), with ultraviolet-visible (UV/Vis) detection system equipped with ULTRA C18 column (5 μ m; 4.6 mm x 250 mm) operating in reverse phase. The mobile phase employed a solution of acidified water with 0.1% acetic acid and acetonitrile with volumetric ratio of 65:35 and a flow of 0.700 mL \cdot min⁻¹. The oven temperature was maintained at 40 \pm 1 $^{\circ}$ C and the pressure at 53 kgf \cdot cm⁻². Then, analytical curves were constructed in a concentration range between 0.1 and 1 mg \cdot L⁻¹. The stock solution (10 mg \cdot L⁻¹) of each pharmaceutical was prepared in a 9:1 mixture of acetonitrile and methanol.

Then, the methodology used was validated and the following parameters were analyzed: linearity, detection limit (LOD), quantification limit (LOQ), precision and accuracy. The linearity evaluation was performed based on the construction of the analytical curve and determination of the correlation coefficient (r). The precision of the method, based on the coefficient of variation (CV), as well as the accuracy were determined based on the recovery method, through the sample fortification procedure. The verification of the quantification and detection limits were performed according to INMETRO (2011). Equations 1 to 4 were used in the determination of such parameters and are set out in Table 1.

Table 1 - Equations used in the validation of the methodology

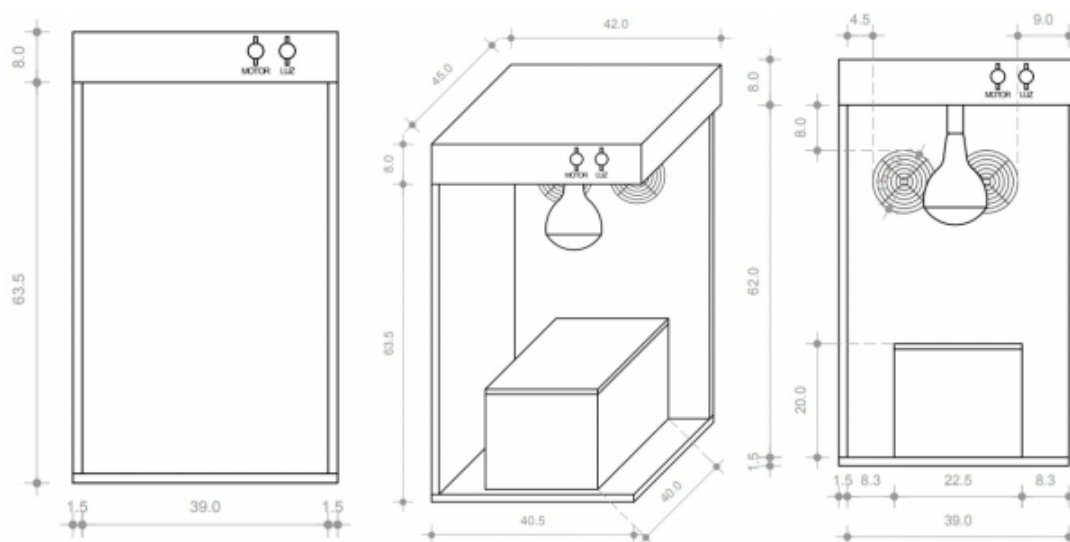
Parameters	Variables	Equation	Number
Precision	X _m it's the arithmetic average of the number of measurements, x _i the individual value of the measurement and n the number of measurements.	$s = (\sum (X_i - X_m)^2 / (n-1))^{1/2}$	(1)
		$CV(\%) = (s / X_m) \cdot 100$	(2)
Accuracy	C ₁ , C ₂ and C ₃ are the concentrations: 1) determined in the sample with the addition of the standard, 2) without addition of the standard and 3) of the standard added.	$R(\%) = ((C_1 - C_2) / C_3) \cdot 100$	(3)
LOQ	s is the estimate of the standard deviation and S the slope of the analytical curve.	$LOQ = 10 \cdot (s/S)$	(4)
LOD		$LOD = 3.3 \cdot (s/S)$	(5)

2.2 Degradation of the pharmaceuticals by AOP using bench reactors

Degradation tests were performed using two types of advanced oxidative processes: photo-Fenton (tests containing iron) and UV/H₂O₂ action (tests in absence of iron). For this, glass beakers (80 mL) filled with 50 mL of an aqueous solution containing the pharmaceuticals under study were used.

Which were placed in a sunlight reactor, with 2 exhaust fans, operating with a 300W ultra-vilatux lamp of the OSRAM brand (Figure 1).

Figure 1 – Schematic representation of the bench reactor using sunlight



Source: Santana et al (2017)

The degradation of the drugs and the best working conditions were evaluated through a factorial design 2^3 with central point (triplicate analysis). The independent variables studied were time, concentration of ($[H_2O_2]$) (F. Maia) and concentration of iron ions ($[Fe]$) ($FeSO_4 \cdot 7H_2O$ – Vetec, being utilized as the iron source). Table 2 contains the planning matrix used in this work, as well as the description of factor levels. It will be seen from this Table that the tests 1 to 4 refer to the photoperoxidation process, whereas the tests 5 to 11 employ the photo-Fenton process.

Table 2 - Factorial design 2^3 + central point (triplicate analysis)

Test	Time (h)	$[H_2O_2]$ ($mg \cdot L^{-1}$)	$[Fe]$ ($mg \cdot L^{-1}$)
1	- (1)	- (14)	- (absence of iron)
2	+ (3)	- (14)	- (absence of iron)
3	- (1)	+ (28)	- (absence of iron)
4	+ (3)	+ (28)	- (absence of iron)
5	- (1)	- (14)	+ (4.4)
6	+ (3)	- (14)	+ (4.4)
7	- (1)	+ (28)	+ (4.4)
8	+ (3)	+ (28)	+ (4.4)
9	0 (2)	0 (21)	0 (2.2)
10	0 (2)	0 (21)	0 (2.2)
11	0 (2)	0 (21)	0 (2.2)

The identification of the working condition was performed through the analysis of total organic carbon (TOC) and chemical oxygen demand (COD). The first of the analyzes was quantified on a Shimadzu high sensitivity TOC equipment (TOC-Vcsh model), with a concentration range of 4 ng·L⁻¹ a 25.000 mg·L⁻¹. The determination of TOC was done indirectly, from the difference between total carbon (TC) and total inorganic carbon (TIC), while the COD measurements were performed using the 5220D spectrophotometric method contained in the Standard methods for the examination of water and wastewater (APHA, 2012).

After the best working condition was determined, the degradation of the drugs via HPLC was evaluated. For this analysis, the solutions underwent an extraction step employing strata-X polymer cartridges operating in reverse phase (500 mg/6 mL - Allcrom). The stationary phase was conditioned with two aliquots of 5 mL of acetonitrile (J.T.Baker), and two aliquots of 5 mL of ultra-pure water.

2.3 Kinetic study: adequacy to the grouped kinetic model

In the best working conditions, identified in the factorial design study, tests were carried out to quantify the kinetic evolution of TOC as a function of time. The suitability of the proposed study to the lumped kinetic model (LKM) was evaluated (ZHANG; CHUANG, 1999). This model describes the profile of the total residual concentration in terms of carbon contained in the liquid phase (Cr), dividing the species formed into three: A) encompasses the pharmaceuticals and intermediates susceptible to oxidation (non-refractory), B) represents the refractory organic species arising from the oxidation of A and C) involves all the carbon dioxide formed from the complete oxidation of organic species.

2.4 Mathematical evaluation: application of artificial neural network

In the artificial neural network was simulated to predict the conversion of TOC, using as input parameters, time, H₂O₂ concentration, iron ions concentration and chemical oxygen demand analyzes. In the ANN construction, to approximate the forward transmission characteristics, a multilayer perceptron (MLP) type configuration was used. For this, the propagation of the network input signal was performed through the neurons distributed in it, as well as in the hidden layer and its output. The MLP used the learning of the backpropagation algorithm of the network, guaranteeing a greater convergence in its learning.

2.5 Toxicity assessment

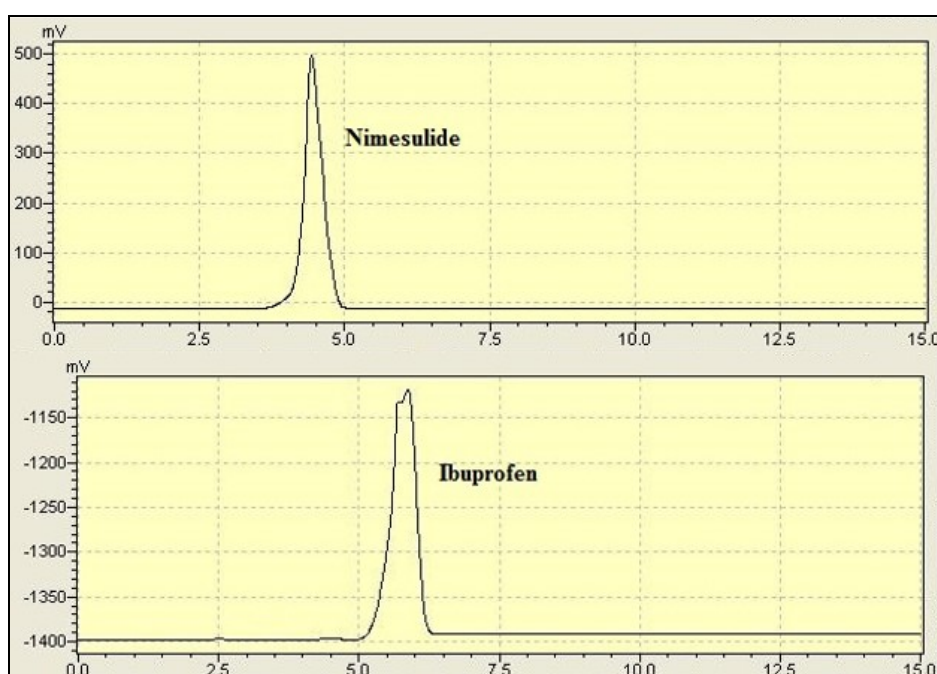
Knowing that after the use of the AOP, intermediates are formed, the toxicity of the treated solution, as well as the samples before treatment, were evaluated using the seeds of *Lactuca Sativa* (lettuce), *Cichorium endívia* (chicory), *Ocimum basilicum* (basil) and *Americano Hard* grain (wheat). For the bioassays were used Petri dishes filled with filter paper, in which 10 seeds and 2 mL of the test solution were placed in each plate. Deionized water was used as negative control and a 3% boric acid solution as positive control. The Petri dishes were kept at 25±1°C, in the absence of light, for a period of 72h (grain) and 120h (seeds). Finally, after the incubation time, the number of germinated seeds and the root growth of each one was observed. Then, the relative growth rate (RGR) and germination index (GI) were calculated according to the methodology proposed by Young et al, 2012.

3 RESULTS AND DISCUSSION

3.1 Identification, quantification and validation of the methodology of the drugs nimesulide and ibuprofen

The pharmaceuticals nimesulide and ibuprofen were analyzed together and observed by HPLC at the wavelengths of 264 and 223 nm, respectively (Figure 3). Such compounds were identified in a retention time range of 3.5 to 6.0 min, which agrees with Maltese; Maugeri; Bucolo (2004) who observed nimesulide in 5.9 min and with Eraga et al (2015) who observed ibuprofen in a time of 3.6 min. Once the suitability of the proposed methodology to identify the drugs under study was verified, the validation step was carried out to guarantee the reliability of the quantification process.

Figure 3 – Chromatograms of the drugs nimesulide and ibuprofen



It is noteworthy that the determination of the nimesulide drug is carried out on the ultraviolet scale since this pharmaceutical has in its molecular structure a weakly acidic methanesulfonamide group, which dissociates the pH dependence of the reaction medium. Pereira et al (2011) also stated that when performing a determination of pKa of nimesulide by spectrophotometry, at pH values between 2 and 10, the presence of isobestic peaks around 270 nm and 340 nm is observed, ensuring that the species can be evaluated with the same safety in both. As for ibuprofen, Borges; Goraieb; Collins (2012) states that at pH between 5 and 7 there are no major changes in the pKa value, so the range used in this work does not compromise the spectrophotometric analysis.

From the dilutions of the stock solution, the analytical curves were constructed, taking as response the areas of the different concentrations analyzed. To evaluate the dispersion of the values of these concentrations, a statistical analysis was performed, based on the calculation of the standard deviation between them, using Grubbs test for 95% confidence (GRUBBS; BECK, 1972).

Table 3 - Average of the areas, standard deviation and Grubbs test for the pharmaceuticals under study

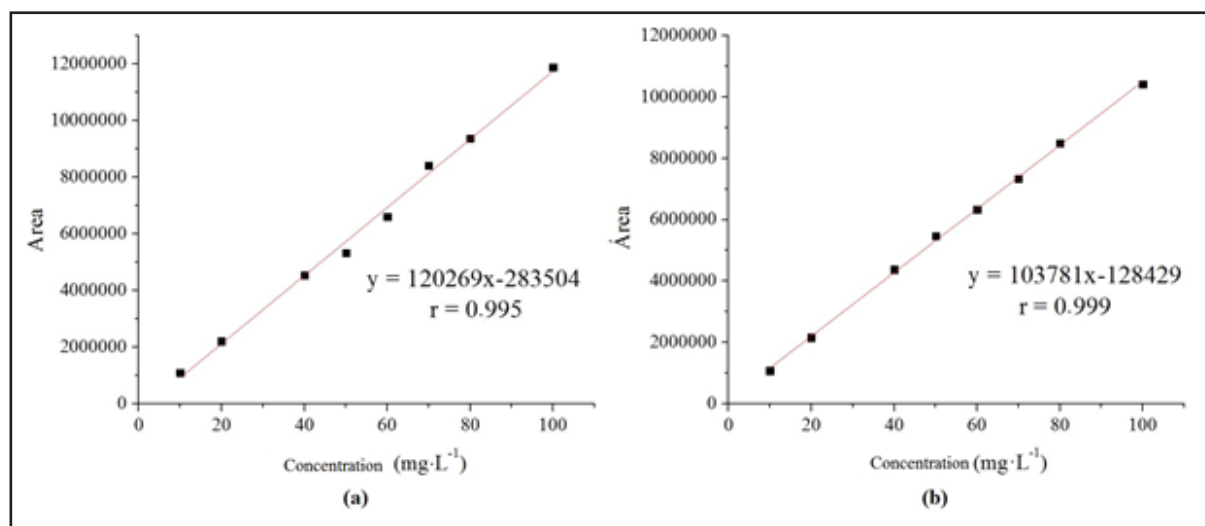
Concentration (mg·L ⁻¹)	Nimesulide				Ibuprofen			
	Average of the areas	Standard deviation	Grubbs Test		Average of the areas	Standard deviation	Grubbs test	
			G _{<}	G _{>}			G _{<}	G _{>}
10	1102602	46589	1.311	1.630	1075693	16769	1.180	1.729
20	2208718	116505	1.275	1.314	2157020	37119	1.260	1.342
40	4541918	95301	1.331	1.243	4380899	80476	1.818	1.312
50	5329161	393931	1.938	0.913	5471956	49130	1.014	1.829
60	6598530	588214	1.605	1.174	6327920	70675	1.617	1.250
70	8418863	207781	1.096	1.445	7328667	92943	1.470	1.540
80	9366057	594617	1.219	1.172	8488089	158593	1.455	1.255
100	11881879	215707	1.758	0.795	10423081	132415	1.473	1.161

*Manual integration method was used to determine peak areas.

Based on the results obtained and described in Table 3, it was possible to verify that the obtained values agreed with each other in the Grubbs test for 97.5% of confidence. It should be noted that for eight measurements and the confidence level chosen, the values of G_< and G_> should be less than 2.126 (GRUBBS; BECK, 1972).

Since the data obtained were coherent, the analysis of linearity was done by constructing the analytical curve and calculating the correlation coefficient (r), as can be observed in Figure 4. It was verified that the analytical curves presented r values of 0.9955 (nimesulide) and 0.9991 (ibuprofen), obeying the standards required by the National Health Surveillance Agency (ANVISA), which considers linear when $r \geq 0.99$, while the National Institute of Metrology, Standardization and Industrial Quality (INMETRO) accepts $r \geq 0.90$ (INMETRO, 2011; BRASIL, 2011).

Figure 4 – Analytical curves of the pharmaceuticals: a) nimesulide e b) ibuprofen



After verifying the linearity of the method for the two pharmaceuticals, the precision was analyzed, based on the quantification of the coefficient of variation. The CV values obtained for each one of the concentrations are described in Table 4.

Table 4 - Values obtained for the coefficient of variation

Concentration (mg·L ⁻¹)	Nimesulide	Ibuprofen	Concentration (mg·L ⁻¹)	Nimesulide	Ibuprofen
10	4.22	1.56	60	8.91	1.12
20	5.27	1.72	70	2.47	1.27
40	2.10	1.84	80	6.35	1.87
50	7.39	0.90	100	1.81	1.27

In order for the method to be considered precise, it is necessary that the values of CV are lower than 20%, which was observed for all concentrations analyzed (INMETRO, 2011). Then the analysis of the accuracy of the method was performed, and the following percentages of recuperation were verified: 97, 103 and 97 for nimesulide and 93, 103 and 118 for ibuprofen. Therefore, an average of 99.0 ± 3.5 for nimesulide and 104.7 ± 12.6 for ibuprofen was obtained, which confirms that the analytical method is accurate, since the recovery percentages obtained are between 50 and 120% (SOUZA, 2011; PERLATTI et al, 2012). Finally, the quantification and detection limits of the methods were determined: a) nimesulide LOQ = 11.83 mg·L⁻¹ and LOD = 3.58 mg·L⁻¹ e b) ibuprofen LOQ = 16.15 mg·L⁻¹ and LOD = 5.33 mg·L⁻¹.

3.2 Degradation of pharmaceuticals by AOP using bench reactor

Initially, for an aqueous solution containing the drug mixture nimesulide (50 mg·L⁻¹) and ibuprofen (50 mg·L⁻¹) the efficiency of the direct photolysis process was evaluated and an organic matter conversion on less than 40% was verified for both compounds. Subsequently, this same solution was subjected to COD analysis, as well as total organic carbon before and after the degradation process. In order to verify the best working condition, a study of factorial planning was carried out, the results which are show in Table 5.

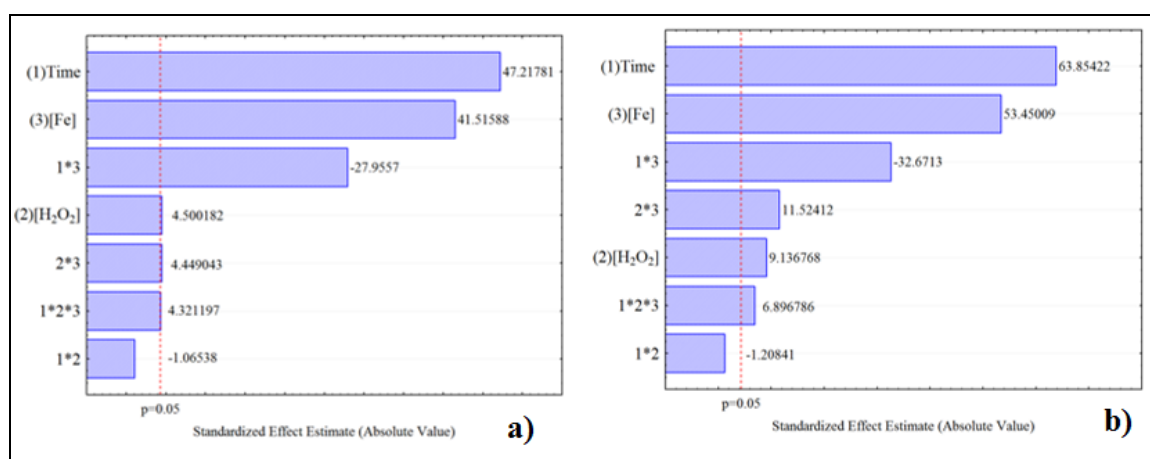
 Table 5 - Results of the tests carried out for factorial design 2³ for the conversion of organic matter (COD and TOC)

Test	TOC (mg de O ₂ ·L ⁻¹)	% of conversion of COD	TOC (mgC·L ⁻¹)	% de conversion of COD
1	29120.0	34.37	1258.0	44.43
2	8154.0	81.63	454.4	79.93
3	27704.0	37.56	1214.0	46.37
4	9537.0	78.50	534.8	76.37
5	11070.0	75.05	632.0	72.08
6	6904.0	84.44	436.2	80.73
7	9587.0	78.39	575.8	77.16
8	3729.0	91.60	233.8	89.67
9	9770.0	77.98	579.2	74.41
10	9070.0	79.55	558.3	75.34
11	9436.0	78.30	573.4	74.67

The analysis of Table 5 allows to verify that the test that presented the best result was number 8, where 91.60% of COD conversion and 89.67% of TOC were observed, this experiment was carried out under the following conditions: time of 3 hours; $[H_2O_2] = 0.12 \text{ g}\cdot\text{L}^{-1}$ e $[Fe] = 0.43 \text{ g}\cdot\text{L}^{-1}$. Thus, for a better evaluation of the data, a statistical analysis was performed by calculating the effects of the factors and their interactions with the Statistica 6.0 program. These identified which of the effects were statistically significant for 95% confidence by constructing Pareto charts (Figure 5).

It is worth noting that, in order to verify that there was no interference of the hydroxyl radicals in the COD analysis, residual peroxide tests were performed using a colorimetric method with Mquant test strips (Merck), with H_2O_2 ranging from 0 to $25 \text{ mg}\cdot\text{L}^{-1}$. It was then verified that the amount of H_2O_2 residual at the end of the treatment was less than $1 \text{ mg}\cdot\text{L}^{-1}$, and the analysis of the organic matter could be performed without interference from the reagents. Similarly, the residual iron concentration presented was properly precipitated with NaOH (0,1 M) by raising the reaction pH to the basic medium so as not to interfere with the TOC analysis.

Figure 5 - Pareto charts referring to the degradation of the pharmaceuticals a) COD conversion e b) TOC conversion

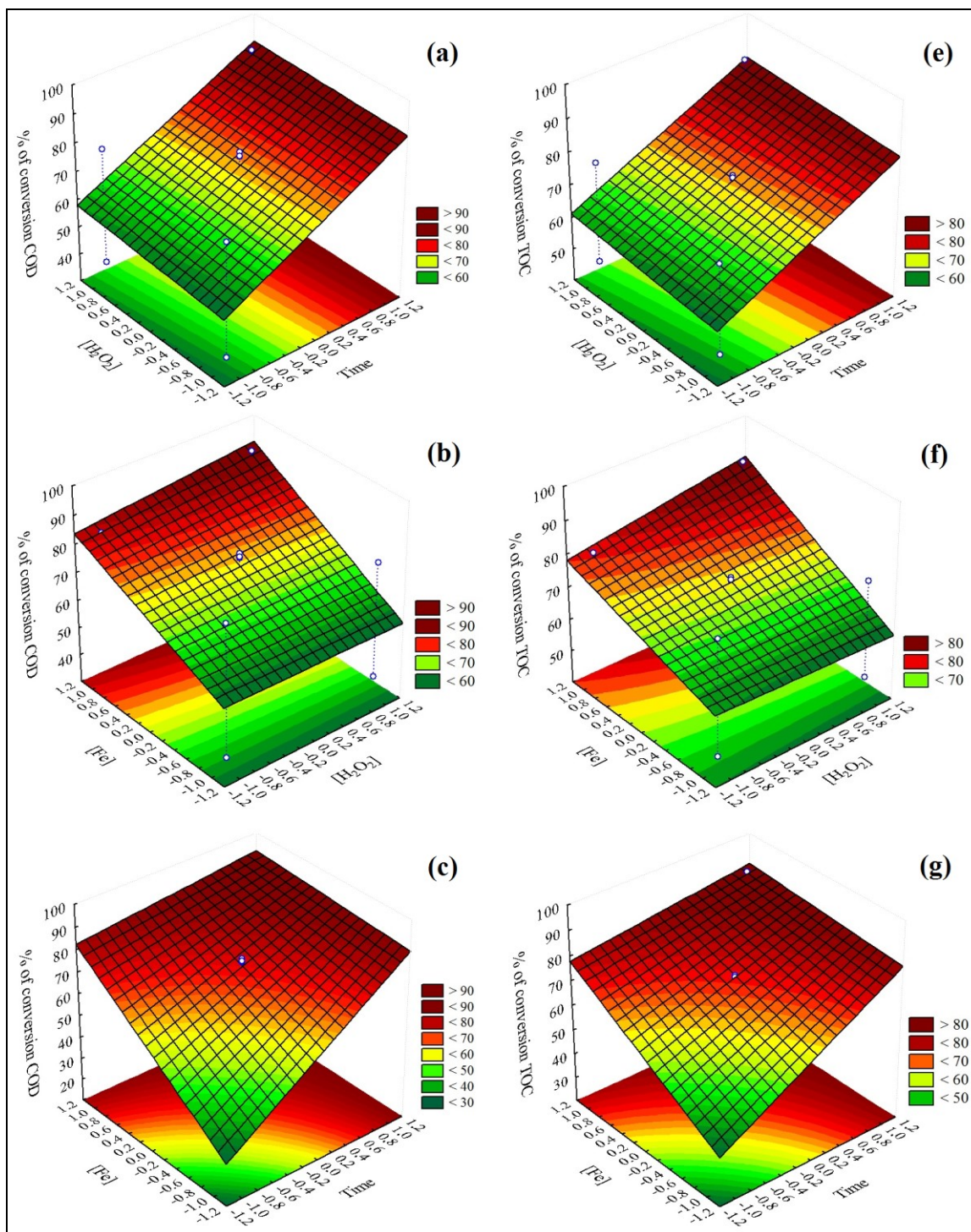


The analysis of figures 5a) and 5b) indicates that all major effects were statistically significant at 95% confidence as well as interactions effects, except for the interaction between $[H_2O_2]$ and time. Thus, for a better understanding of the behavior of the interactions between the variables studied in the conversion of COD and TOC, surface graphs were generated, which can be observed in Figure 6.

The analysis of Figures 6a) and 6e) indicates that the interaction of time and $[H_2O_2]$ variables does not show large variations in the values of organic matter conversion, confirming that observed in the Pareto charts. (Figure 5). When analyzing Figures 6b) and 6f) it is verified that when the highest levels of $[Fe]$ and $[H_2O_2]$ are combined, a greater conversion of COD and TOC is obtained. Finally, when analyzing the combination between $[Fe]$ and time, it is observed that a higher conversion is observed for the combination between the higher levels of the two analyzed variables (Figures 6c) and 6g)). Thus, it can be stated that the working conditions obtained in the present study, regarding the iron concentration value, meet the water release standards established by the resolution of National Environmental Council (CONAMA) number 430/2011, indicating that the $[Fe]$ should be less than $15 \text{ mg}\cdot\text{L}^{-1}$ (CONAMA, 2011).

Determined the best process conditions ($[H_2O_2] = 28.0 \text{ mg}\cdot\text{L}^{-1}$ and $[Fe] = 4.4 \text{ mg}\cdot\text{L}^{-1}$), a kinetic study of the pharmaceuticals degradation using the photo-Fenton process was carried out. For this study, the total organic carbon concentrations were used as response to verify the adequacy to the grouped kinetic model, as well as to evaluate the application of artificial neural networks to predict a mathematical model.

Figure 6 - Response Surface of the interaction: a) and e) $[H_2O_2]$ and time; b) and f) $[Fe]$ and $[H_2O_2]$; c) and g) $[Fe]$ and time



3.3 Kinetic study: adequacy to the grouped kinetic model

In the best working condition, the concentrations of the two drugs under study were evaluated by HPLC, finding that the photo-Fenton process degraded 89.70% of nimesulide and 93.35% of ibuprofen. Based on the best experimental conditions described above, a kinetic study was carried out by monitoring the total organic carbon concentration. The description of the experimental results for the TOC concentration together with the relation between the TOC and the initial TOC (TOC_0) are described in Table 6.

Table 6 - Evolution of TOC concentration over time and the ratio of TOC to the initial TOC (TOC_0), observed and calculated

Time (min)	TOC (mgC·L ⁻¹)	TOC/ TOC_0 (observed)	TOC/ TOC_0 (calculated)
0	6053.00	1.00	1.00
15	5462.00	0.90	0.91
30	4855.00	0.80	0.82
60	4003.00	0.66	0.63
120	2148.00	0.35	0.35
180	921.50	0.15	0.19
240	608.70	0.10	0.10
360	602.80	0.10	0.03

By analyzing Table 6 it is possible to verify that a conversion of 90.04% of the organic matter was reached after 360 min ($[COT] = 602.80 \text{ mgC}\cdot\text{L}^{-1}$). With these data, we could then verify the adequacy to the kinetic group model as well as to determine the apparent velocity constants k_1 , k_2 , e k_3 from Equation 8. The values of the constants were obtained through non-linear interpolation applied to the data found using the *Microsoft Office Excel* solver tool. The method of minimizing the quadratic sum of the residues was used in the software.

$$(TOC/TOC_0) = C_r = [(k_1 - k_3) / (k_1 + k_2 - k_3)] e^{-(k_1+k_2)t} + [k_2/(k_1 + k_2 - k_3)] e^{-k_3t} \quad (8)$$

Thus, based on Equation 8, the values of k_1 , k_2 e k_3 were determined being respectively 0.024 min^{-1} , 0.010 min^{-1} e 0.005 min^{-1} . These results indicate that the mineralization of the pharmaceuticals plus the non-refractory intermediates ($k_1 = 0.024 \text{ min}^{-1}$) occurs at a higher reaction rate than the degradation in refractory intermediates ($k_2 = 0.010 \text{ min}^{-1}$). However, the mineralization of the refractory intermediates has a lower reaction rate ($k_3 = 0.005 \text{ min}^{-1}$) than the other reactions involved. This indicates that the group of pharmaceuticals plus the non-refractory intermediates has a greater tendency of mineralization when compared to the formation of refractory intermediates. Thus, the graphs for the adjusted kinetic model fit and the comparative analysis of the theoretical and experimental TOC values can be constructed, as can be observed in Figure 7.

Figure 7 - a) Adjustment of the kinetic model grouped to the experimental data of the TOC conversion; b) Comparison between theoretical and experimental TOC values

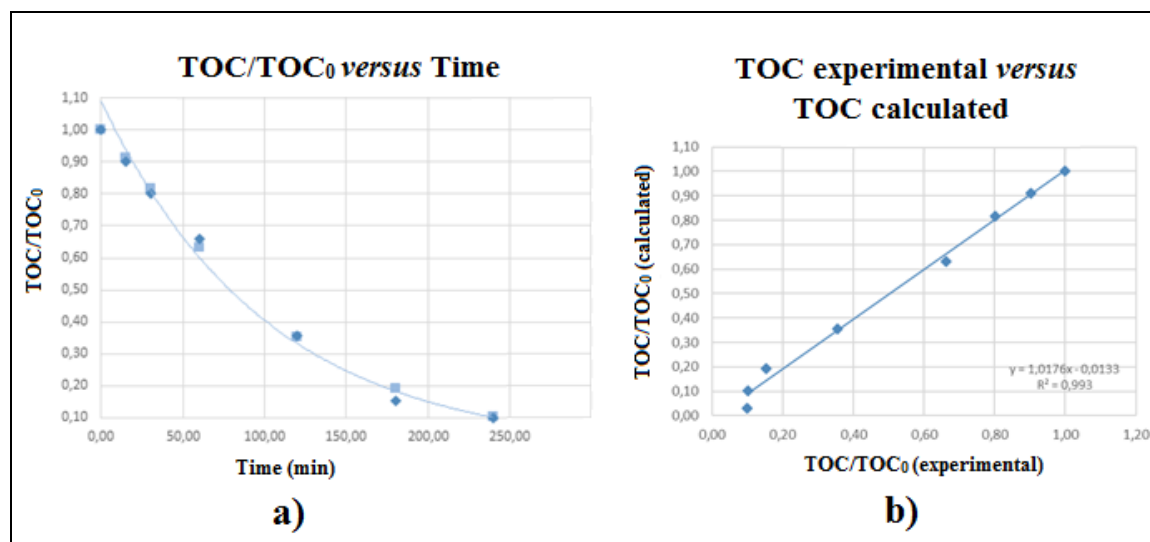


Figure 7 show a good correlation between the data obtained experimentally and the calculated theoretical data, indicating that the adopted kinetic model can be applied satisfactorily to the TOC conversion over time. This verification is corroborated by the value of the linear regression coefficient (R^2) that was equal to 0.993.

3.4 Mathematical evaluation: application of artificial neural networks

Once the kinetic data for degradation of the pharmaceuticals ibuprofen and nimesulide were found to be well adapted to the grouped kinetic model, the mathematical evaluation using artificial neural networks was applied. The descriptive statistics of the five variables used in ANN are contained in Table 7.

Table 7 - Descriptive statistics of the five variables used in ANN

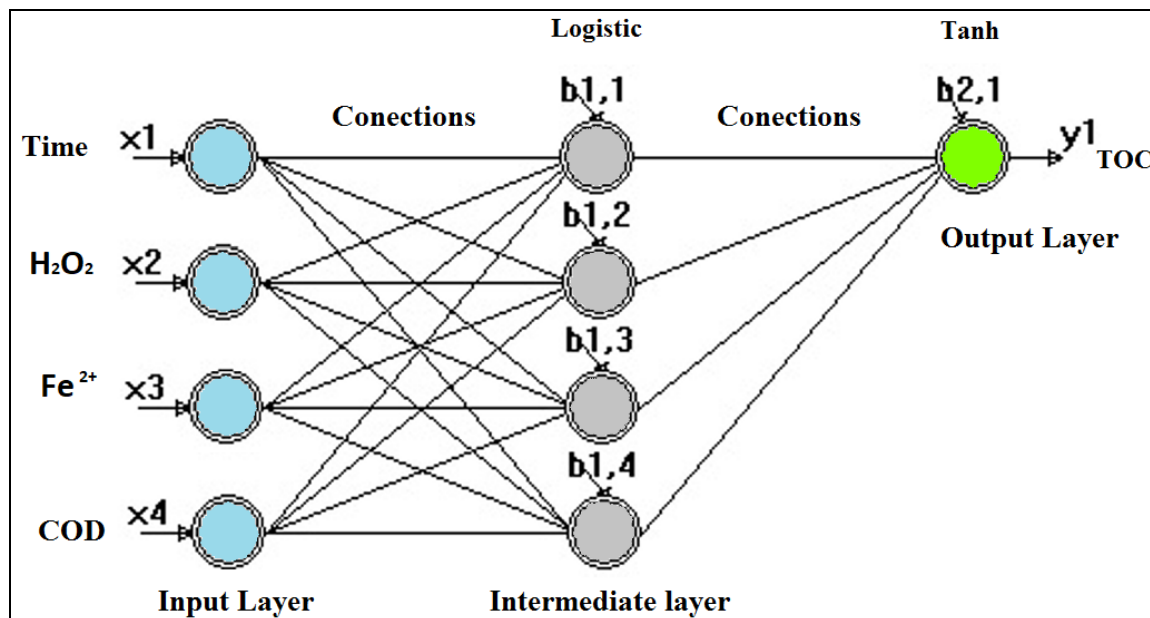
Variables	Training				Test				Validation			
	Min	Max	Average	σ	Min	Max	Average	σ	Min	Max	Average	σ
Tempo (min)	0.00	360.00	122.86	86.28	0.00	360.00	155.00	129.03	0.00	300.00	146.67	106.49
[H ₂ O ₂] (mg·L ⁻¹)	14.00	28.00	23.67	5.77	28.00	28.00	28.00	0.00	14.00	28.00	22.56	6.65
[Fe ²⁺] (mg·L ⁻¹)	0.00	4.40	2.93	1.86	0.00	4.40	3.42	1.94	2.20	4.40	4.16	2.09
COD(%)	0.00	91.60	65.12	23.95	0.00	86.82	55.27	30.87	0.00	85.73	67.73	36.31
TOC(%)	0.00	90.04	63.08	26.07	0.00	90.04	60.23	36.09	0.00	89.99	68.65	40.26

Min = minimum, Max = maximum, σ = standard deviation

Through the analysis of Table 7 it is possible to verify in the training and in the test a conversion of 90.04%, while in the validation the conversion rate of total organic carbon was 89.99%. The optimization of the ANN topology is probably the most important step in the development of the

model. In the present work, a three-layer posterior propagation neural network (4:4:1) was used to model organic matter conversion. Figure 8 shows the architecture diagram ANN MLP 4-4-1 BFGS 4567.

Figure 8 - Architecture diagram ANN MLP 4-4-1 BFGS 4567



Thus, the ANN for the photo-Fenton/sunlight system used in the degradation of nimesulide and ibuprofen pharmaceuticals presented a linear regression coefficient (R^2) equal to 1.00 for training, testing and validation when using MLP model 4-1-1. The training algorithm used was the BFGS 4567, the SOS error function, using as an activation function an internal layer logistic and output Tanh.

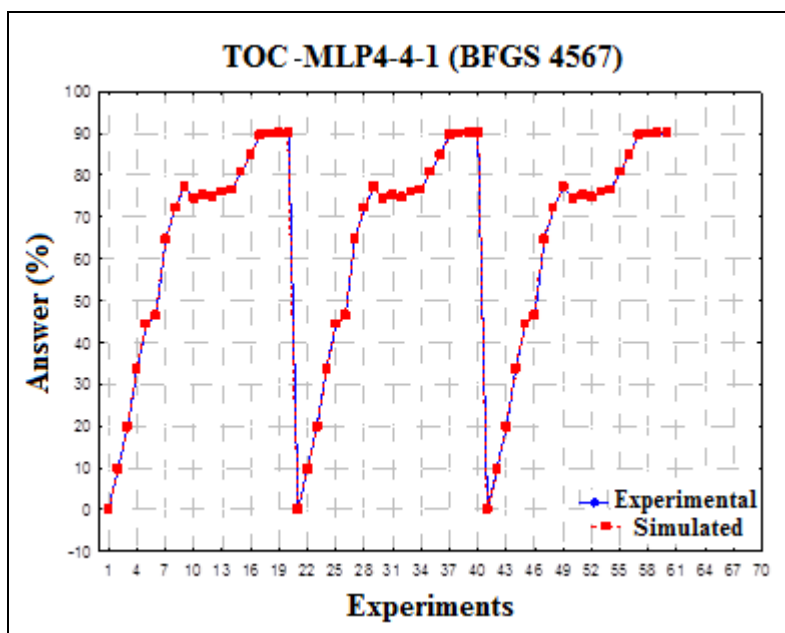
To calculate training, validation and test errors, all outputs were performed on an inverse interval scale to return the predicted responses to their original scale. Then, these data were compared with the values obtained experimentally. The ANN used in this work provided the weights listed in Table 8.

Table 8 - Values of the weights of the entry, intermediates and exit layers of ANN MLP 4-4-1 (BFGS 4567, Logistic - Tanh)

Neuron	Weights					
	Time (min)	[H ₂ O ₂] (mg.L ⁻¹)	[Fe ²⁺] (mg.L ⁻¹)	COD (%)	Bias input	TOC conversion (%)
1	1.3351	2.4592	6.9613	2.7625	-15.8208	66.5294
2	-3.1948	-34.9400	-35.4083	36.8891	7.7467	1.3869
3	29.3356	12.3996	-4.7303	15.4220	-6.4942	46.0932
4	-30.9336	-24.3306	2.1527	27.7620	21.0440	-14.4343
Bias intermediate						-31.8136

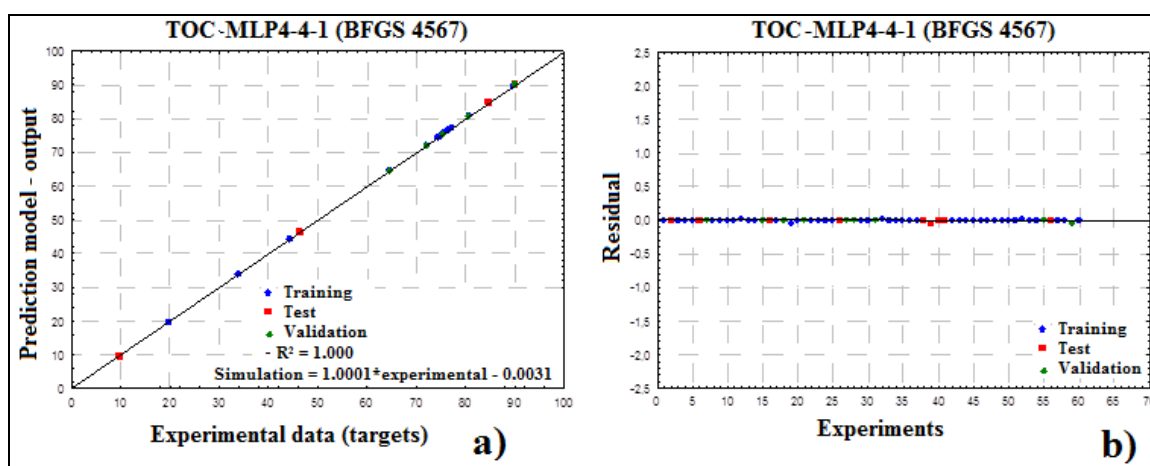
Figure 8 shows the graph comparison between the experimental results and the simulated values by the MLP network (BFGS 4567). The analysis of this Figure allows to verify a good adequacy of the experimental results, being able to affirm that the ANN used is accurate.

Figure 9 - Comparison between experimental results and simulated MLP (BFGS 4567)



After analyzing Figure 9 and verifying the fit of the experimental data to the simulated values by the MLP network (BFGS 4567), the linear and residual regression graphs were generated (Figure 10). In Figure 10a) it can be verified that the prediction model presented a good fit, with a linear regression coefficient (R^2) equal to 1.000, confirming that the neural network model reproduces the conversion of the TOC in the system, within the ranges used in the assembly model.

Figure 10 - a) Linear regression between the experimental and simulated results of MLP ANN (BFGS 4567), b) Residues of the conversion of experimental TOC and simulated by ANN MLP (BFGS 4567)



It can be stated, therefore, that the ANN model employed represents satisfactorily the behavior of the degradation process within the analyzed domain. This fact can be confirmed by the graph of residues from the conversion of the experimental TOC and simulated by MLP ANN (BFGS 4567) (Figure 10b). Through this Figure, the data of the residual analysis did not show dispersion tendencies around the x-axis.

3.5 Toxicity analysis

To evaluate the toxic effects of the aqueous solution before and after the AOP treatment, toxicity tests were performed. The results of the bioassay averages, in triplicate, are shown in Table 9. It is worth mentioning that the tests performed with the positive control did not present germination.

Table 9 - Average germinated seed for each species studied

Sample	<i>Americano hard</i>	<i>Cichorium endívia</i>	<i>Lactuta Sativa</i>	<i>Ocimum basilicum</i>
Water	9.00 ± 0.00	8.33 ± 0.58	8.33 ± 0.58	8.66 ± 0.58
SBD	5.33 ± 0.58	1.00 ± 1.00	5.66 ± 0.58	3.33 ± 0.58
SAD	9.33 ± 0.58	9.33 ± 0.58	8.66 ± 1.53	9.66 ± 0.58

(*)SAD = solution before degradation; SAD = solution after degradation

From Table 9 it can be inferred that number of seeds germinated when in contact with the solution before treatment were less when compared with the ones in contact with water, which indicates the presence of toxicity. A similar result was obtained by researchers who evaluated the toxicity of pharmaceutical effluent for seeds, in which an inhibition of germination was verified (NAPOLEÃO et al, 2018).

The solution after the treatment behaved analogously to the negative control for the different concentrations evaluated, being possible to state that the compounds formed after the applied treatment do not present toxicity in relation to the species studied, with respect to germination. However, this study is not sufficient to assert that there is no toxicity of the analyzed compounds, and it is necessary to verify the root length (Table 10).

Table 10 - Average root growth for each species studied

Sample	<i>Americano hard</i>	<i>Cichorium endívia</i>	<i>Lactuta sativa</i>	<i>Ocimum basilicum</i>
Water	7.20 ± 0.75	2.20 ± 0.10	5.83 ± 0.60	2.23 ± 0.25
SBD	4.10 ± 0.17	0.53 ± 0.31	1.47 ± 0.59	0.53 ± 0.45
SAD	7.37 ± 0.60	2.27 ± 0.38	5.23 ± 0.64	2.47 ± 0.15

(*)SBD = solution before degradation; SAD = solution after degradation.

The results shown in Table 10 show that although some seeds may survive in potentially toxic medium (SBD), their development has suffered sublethal effects (inhibition of root development). Whereas after the treatment, no relative toxic effect was observed in the seeds. These data agree with those obtained by Napoleão et al (2014), who evaluated the toxicity of a pharmaceutical effluent submitted to the photo-Fenton process, against the *Lactuta sativa* seed and the *Americano Hard* grain. Finally, for a better understanding of the results, the germination index (GI) and the relative growth rate (RGR) were determined for the studied species, as can be observed in Table 11.

Table 11 - RGR and GI values (%) for each studied species

Sample	<i>Americano hard</i>		<i>Cichorium endívia</i>		<i>Lactuta sativa</i>		<i>Ocimum basilicum</i>	
	RGR	GI (%)	RGR	GI (%)	RGR	GI (%)	RGR	GI (%)
Water	1.00	100.00	1.00	100.00	1.00	100.00	1.00	100.00
SBD	0.57	33.72	0.24	2.91	0.25	17.08	0.24	9.18
SAD	1.02	106.07	1.03	115.40	0.90	93.27	1.10	123.20

(*)SBD = solution before degradation; SAD = solution after degradation

The analysis of Table 11 shows that there was interference in both the RGR and the GI of all studied species in the solution before treatment. On the other hand, the solution treated through the photo-Fenton process did not show a significant difference in relation to the indices evaluated, when compared to the behavior of the negative control. This indicates that the solution after treatment is less toxic than the initial sample containing the pharmaceuticals nimesulide and ibuprofen.

4 CONCLUSION

The present work allowed to verify that the photo-Fenton process was efficient in the treatment of the drugs nimesulide and ibuprofen, degrading 89.70% and 93.35% respectively. Besides reducing the concentration of chemical oxygen demand by 91.60% and mineralizing 90.04% of the organic matter by reducing the levels of total organic carbon. It was verified that the kinetic study presented a good linear fit, with R^2 equal to 0.993 for the clustered kinetic model, as well as the artificial neural network MLP 4-4-1 BFGS 4567 can predict satisfactorily the data for the proposed treatment, having as response the TOC concentrations. Finally, the toxicity of the solution after the photo-Fenton process against the seeds of *Lactuta Sativa* (lettuce), *Cichorium endívia* (chicory), *Ocimum basilicum* (basil) and *Americano Hard* grain (wheat) were analyzed. Verifying that the seeds that received the solution before the AOP had a reduction of the germination, which was not observed when adding the solution after treatment. To complete the toxicity study, the root growth of the studied species was evaluated and no toxic effect was detected.

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