INSTRUCCIONES PARA LOS AUTORES

El archivo debe ser un documento de Word, su extensión deberá ser doc o docx. El archivo no podrá pesar más de 2 Megabytes.

Las reseñas de Tesis Doctorales se publicarán en inglés.

La extensión para cada reseña es de 4 páginas máximo (Times New Roman 11, interlineado 1.5)

Las reseñas constarán de los siguientes apartados:

1. Título de la Tesis Doctoral.
2. Nombre y apellido del doctorando, año de lectura de la Tesis Doctoral y centro o empresa donde ha sido realizada.
3. Director/es de la Tesis Doctoral y su afiliación.
4. Redacción del texto y presentación. Se procurará que la redacción sea lo más clara y concisa posible, incluyendo: Objectives and Novelty, Results, Conclusions and Related Publications.

Se debe dar especial énfasis a los objetivos y novedad de la Tesis Doctoral, la necesidad de la investigación y las conclusiones obtenidas.

Las Tablas y Figuras deben insertarse en el texto.

En el apartado de Related Publications, la bibliografía deberá reducirse a la que tenga relación directa con la Tesis Doctoral. Las referencias deben presentarse en una lista numérica al final de la reseña de acuerdo al formato que se muestra en las instrucciones para los artículos del Boletín.

Finalmente se debe incluir un link al final de la reseña para descargar la Tesis Doctoral completa en su versión original.

A continuación se muestra una Reseña de Tesis Doctoral, a modo de ejemplo, que sigue estas instrucciones.
DEGRADATION AND REMOVAL OF NAPHTHALENESULPHONIC ACIDS IN AQUEOUS SOLUTION BY MEANS OF OZONATION PROMOTED BY ACTIVATED CARBON

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OBJECTIVES AND NOVELTY
Effluents from the textile industry present elevated concentrations of sulphonated polyphenols (dyestuffs), ranging between 10 and 80 mg/L and with a mean concentration of 31 mg/L. Biological water treatments are not effective in eliminating this type of compound, so that alternative treatments are necessary. At present, activated carbon and ozone are widely used to treat organic compounds that are resistant to biological treatments, taking advantage of their respective adsorbent and oxidizing properties. Activated carbon has been signaled as a promising catalyst of ozonation because of its chemical and textural properties and low cost. However, there are no data on the physical and chemical processes that take place to enable the correct design of a reactor for this method. Thus, it is not known which properties of activated carbon are involved in increasing the extension of the ozonation process or what reactions are implicated. Moreover, the influence of operational variables on the adsorption/ozonation process is still unknown.

The present work aimed to outline results obtained during the development of a wider project to design a new reactor based on the combined use of ozone and activated carbon for the depuration of waters that contain a large concentration of non-biodegradable organic matter. Knowledge of the ozonation process of naphthalenesulphonic acids in the absence of activated carbon (reaction mechanism and kinetics, influence of operational variables, oxidation subproducts) and of the interactions involved in the adsorption of these contaminants on activated carbon will also help us explain more precisely the behavior of activated carbon both as catalyst and adsorbent in the treatment system based on the combined use of ozone and activated carbon.
RESULTS

Naphthalenesulphonic compounds were observed to present genotoxic activity at elevated concentrations. The study revealed that the genotoxic activity of naphthalenesulphonic acids reduces as the number of sulphonic groups on the aromatic ring increases. Ames test findings indicate that the mechanism that produces the DNA damage is by substitution of nitrogenated bases. The aromatic acids studied showed a low reactivity to ozone. The greater the number of sulphonic groups in the molecule, the lower was the reactivity. The prolonged exposure of activated carbon to ozone modifies the chemical composition of the surface of activated carbon. The basic sites are transformed into acid ones due to their oxidation. The action of ozone also impacts on the textural characteristics of the carbon; the surface area diminishes. On the other hand, the adsorption of naphthalenesulphonic acids on activated carbons is regulated both by $\pi$-$\pi$ dispersion interactions and by electrostatic interactions. We reported that the reactivity of naphthalenetrisulfphonic acid (NTS) to ozone was very low ($k_D = 6.72 \text{ M}^{-1} \text{s}^{-1}$). However, NTS showed an elevated affinity for free radicals ($k_{OH} = 3.68 \times 10^9 \text{ M}^{-1} \text{s}^{-1}$), so that we can consider NTS an ideal compound to detect the presence of highly oxidizing radicals in the medium.

To determine the properties of activated carbons involved in the decomposition of ozone in highly oxidizing species, NTS was ozonized in the presence of different commercial activated carbons. Figure 1 shows experimental results of NTS ozonation in the presence of the different activated carbons.

Figure. 1. NTS ozonation in the presence of commercial activated carbons. pH 2.3, T 298 K. (◇), without carbon; (○), Filtrasorb 400; (△), Merck; (⋄), Ceca GAC; (Θ), Ceca AC40; (x), Norit; (+), Sorbo; (-) Witco.
The greater rate of NTS ozonation in the presence of these carbons could, therefore, be explained by an increase in free radical hydroxyl concentration. The carbons that most enhanced the NTS ozonation rate were those with greatest pH_{pzc} values and highest concentrations of surface basic groups. However, no clear relationship was observed between the NTS ozonation rate and the S_{tot} of the activated carbon. On the other hand, the carbons that most favored the removal of NTS from the medium (Sorbo and Norit) were those with greatest macropore volumes. The results obtained suggest that the catalytic activity of these activated carbons in NTS ozonation is mainly a function of the carbon basicity. The basicity of an activated carbon is due to the presence of basic oxygen-containing functional groups (e.g. pyrones or chromenes) and/or graphene layers acting as Lewis bases and forming electron donor-acceptor (EDA) complexes with H_{2}O molecules. According to the above results, we proposed that the delocalised \( \pi \) electron system of basic carbons and oxygenated basic groups (chromene and pyrone) would, therefore, act as catalytic centres of reaction, reducing the ozone molecules to hydroxyl ion and hydrogen peroxide following the reactions (1, 2), which act as ozone decomposition initiators.

\[
O_3 + H_2O + 2e^- \leftrightarrow O_2 + 2 OH^- \quad (1)
\]

\[
O_2 + O_3 \rightarrow \text{O}_3\text{O}_2 \quad (2)
\]

**CONCLUSIONS**

The aromatic acids studied showed a low reactivity to ozone. The greater the number of sulphonic groups in the molecule, the lower was the reactivity.

The prolonged exposure of activated carbon to ozone modifies the chemical composition of the surface of activated carbon. The basic sites are transformed into acid ones due to their oxidation. The action of ozone also impacts on the textural characteristics of the carbon; the surface area diminishes because of the ozone attack and because the increase in oxygenated groups prevents the diffusion of nitrogen by obstructing the entrances of the micropores.

The adsorption of naphthalenesulphonic acids on activated carbons is regulated both by \( \pi-\pi \) dispersion interactions and by electrostatic interactions. However, the \( \pi-\pi \) interactions have a more determining influence on this process.
The presence of basic activated carbon during the ozonation of NTS increases the rate of degradation and produces the removal of part of the organic matter by its mineralization.

**RELATED PUBLICATIONS**


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