

Book of Abstracts

C XI National Meeting on
Catalysis and Porous Materials

C II Meeting of the
Carbon Group

9th - 10th december 2021, Aveiro, Portugal

Title

Abstract book of the XI National Meeting on Catalysis and Porous Materials and the II Meeting of the Carbon Group

Editors

Diana C. G. A. Pinto, Martyn Pillinger, Anabela Valente and Mário Simões

Photo Editor

José M G Pereira

Treatment of leachate waters by wet peroxide oxidation with a compost-based catalyst: effect of pH

Gabriel F. Batista^{a,b}, Fernanda F. Roman^{a,c}, Jose L. Diaz de Tuesta^a, Raquel V. Mambrini^b,
Helder T. Gomes^a

^aCentro de Investigação de Montanha (CIMO), Instituto Politécnico de Bragança, 5300-253 Bragança, Portugal. ^bDepartamento de Química, Centro Federal de Educação Tecnológica de Minas Gerais - CEFET-MG, 30421-169 Belo Horizonte, MG, Brasil. ^cLaboratory of Separation and Reaction Engineering – Laboratory of Catalysis and Materials (LSRE-LCM), Faculdade de Engenharia, Universidade do Porto, 4200-465 Porto, Portugal. E-mail: htgomes@ipb.pt

A compost-based catalyst was synthesized by hydrothermal carbonization following the procedure described elsewhere¹ (3 g of compost in 30 mL of water, 230 °C for 2 h). The material was assessed in the catalytic wet peroxide oxidation (CWPO) of a leachate water, generated during an anaerobic digestion of municipal solid waste. The leachate water is characterized by a high pollutant load (chemical oxygen demand, COD, of 60 g L⁻¹ and total organic carbon, TOC, of 27 g L⁻¹). The CWPO runs were conducted at initial pH (pH₀) of 3 and 6, and at the natural pH of the effluent (7.2), C_{catalyst} = 1.8 g L⁻¹, 80 °C, and the stoichiometric concentration of H₂O₂ needed to mineralize the organic content (based on COD). Fig. 1 shows the results obtained along the reaction. An acidic pH (pH₀ = 3) resulted in a more controlled, but also incomplete, consumption of H₂O₂, leading to a low conversion of COD and TOC (20 and 10%, respectively). Contrarily, the natural pH led to a very fast and uncontrolled consumption of the oxidant source, resulting in 100% decomposition of H₂O₂ in less than 2 h of reaction, but failing to remove COD or TOC (negligible removal, ca. 0%), ascribed to parasitic reactions occurring by the inefficient consumption of H₂O₂. At pH₀ = 6, an intermediate behavior was observed: complete decomposition of H₂O₂ was possible, at a more controlled rate compared to the natural pH. The result was an increment in COD (41%) and TOC removals (19%), almost two times than that observed at the pH₀ 3.

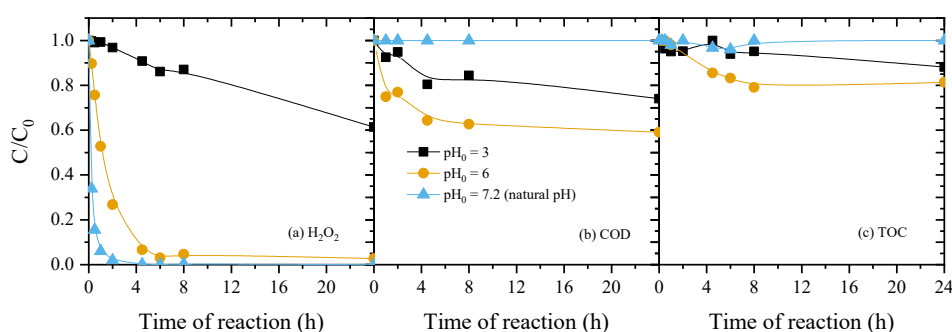


Figure 1. Effect of pH on (a) H₂O₂ concentration, (b) COD removal, and (c) TOC removal.

References

[1] Roman, F. F.; Diaz de Tuesta, J. L.; Praça, P.; Silva, A. M. T.; Faria, J. L.; Gomes, H. T.; *J. Environ. Chem. Eng.*, **2021**, 9, 104888.

Acknowledgments: This work was financially supported by project “VALORCOMP - Valorización de compost y otros desechos procedentes de la fracción orgánica de los residuos municipales”, with reference 0119 VALORCOMP_2_P, through FEDER under Program INTERREG; Base Funding - UIDB/50020/2020 of the Associate Laboratory LSRE-LCM - funded by national funds through FCT/MCTES (PIDDAC); CIMO (UIDB/00690/2020) through FEDER under Program PT2020, and national funding by FCT, Foundation for Science and Technology, and European Social Fund, FSE, through the individual research grant SFRH/BD/143224/2019 of Fernanda Fontana Roman.