

Editorial

Editorial Catalysts: Special Issue on Trends in Catalytic Wet Peroxide Oxidation Processes

Asunción Quintanilla * and Macarena Munoz *

Chemical Engineering Department, Universidad Autónoma de Madrid, Ctra. Colmenar km 15, 28049 Madrid, Spain

* Correspondence: asun.quintanilla@uam.es (A.Q.); macarena.munnoz@uam.es (M.M.); Tel.: +34-91-497-3454 (A.Q. & M.M.); Fax: +34-91497-3516 (A.Q. & M.M.)

Received: 29 October 2019; Accepted: 31 October 2019; Published: 4 November 2019



The catalytic wet peroxide oxidation (CWPO) process is an advanced oxidation technology that has shown great potential for the decontamination of wastewater. CWPO allows the removal of recalcitrant organic compounds under mild conditions (temperatures and pressures in the range of 25–100 °C and 0.1–0.5 MPa, respectively) by using hydrogen peroxide (H₂O₂) as an oxidant, which is considered an environmentally friendly agent. This process requires a solid catalyst with redox properties to generate hydroxyl and hydroperoxyl radicals from the H₂O₂ decomposition. These radical species easily react with the pollutants, oxidizing them into biodegradable forms and finally into CO₂ and water.

This special issue gives an overview of the state-of-the-art CWPO research for the treatment of industrial and urban wastewaters and how this process can be integrated into the water treatment process [1]. It is illustrated that the high versatility of this low-cost technology, thanks to the CWPO operational flexibility, is easily adaptable to any kind of wastewater, either polluted by high-loaded recalcitrant organics in industrial wastewaters or by emerging pollutants at micro-concentration levels in urban waters. This versatility also stands on the application of different types of solid catalysts, which can be tailored according to the process requirements.

For this reason, intensive research effort has been focused on the development of catalysts capable of promoting the abatement of different pollutants in combination with an adequate stability for long-term use and high efficiency of H₂O₂ consumption. In this sense, supported gold nanoparticles have demonstrated to fit these requirements, and a rigorous revision of the main goals of CWPO in presence of gold catalyst can be found in the special issue [2]. However, deactivation cannot be completely avoided due to progressive fouling of the catalyst by the condensation by-products formed upon reaction. An insight into the CWPO reaction mechanism in order to understand the formation, nature, and role of these species [3,4] as well as the hydroxyl radical production mechanism [5], has been also covered.

On the other hand, different innovative solutions show the current trends in the CWPO technology, mainly aimed at the development of an efficient process operated at ambient conditions, by assisting CWPO with UV light irradiation [6], solar light [7], air flow [8], or additional radical activators [9,10]; and also by operated under neutral pH with efficient production of hydroxyl radicals [11]. All these achievements, with significant impact on the operating cost of the CWPO units, were conditioned by the presence of a proper catalyst designed and tailored to provide the best performance.

Finally, we would like to acknowledge the work of excellence developed by the authors of all the contributions to this collection issue, the good aid provided by the involved editorial assistants, and the efforts and comments provided by the reviewers to improve the quality of the articles. Without them, this special issue would not have been possible.

Conflicts of Interest: The authors declare no conflict of interest.

References

1. Rueda Márquez, J.J.; Levchuk, I.; Sillanpää, M. Application of catalytic wet peroxide oxidation for industrial and urban wastewater treatment: A review. *Catalysts* **2018**, *8*, 673. [[CrossRef](#)]
2. Rodriguez, C.S.D.; Silva, R.M.; Carabineiro, S.A.C.; Maldonado-Hódar, F.J.; Madeira, L.M. Wastewater treatment by catalytic wet peroxidation using nano gold-based catalysts: A review. *Catalysts* **2019**, *9*, 478. [[CrossRef](#)]
3. Quintanilla, A.; Díaz de Tuesta, J.L.; Figueruelo, C.; Munoz, M.; Casas, J.A. Condensation by-products in wet peroxide oxidation: Fouling or catalytic promotion? Part I: Evidences of an autocatalytic process. *Catalysts* **2019**, *9*, 516. [[CrossRef](#)]
4. Quintanilla, A.; Díaz de Tuesta, J.L.; Figueruelo, C.; Munoz, M.; Casas, J.A. Condensation by-products in wet peroxide oxidation: Fouling or catalytic promotion? Part II: Activity, nature and stability. *Catalysts* **2019**, *9*, 518. [[CrossRef](#)]
5. Liu, Z.; Shen, Q.; Zhou, C.; Fang, L.; Yang, M.; Xia, T. Kinetic and mechanistic study on catalytic decomposition of hydrogen peroxide on carbon-nanodots/graphitic carbon nitride composite. *Catalysts* **2018**, *8*, 445. [[CrossRef](#)]
6. Sarto, S.; Paesal, P.; Tanyong, I.B.; Laksmana, W.T.; Prasetya, A.; Ariyanto, T. Catalytic degradation of textile wastewater effluent by peroxide oxidation assisted by UV light irradiation. *Catalysts* **2019**, *9*, 509. [[CrossRef](#)]
7. Xu, X.; Liu, S.; Cui, Y.; Wang, X.; Smith, K.; Wang, Y. Solar-driven removal of 1,4-dioxane using WO₃/γ-Al₂O₃ nano-catalyst in water. *Catalysts* **2019**, *9*, 389. [[CrossRef](#)]
8. Campos, A.M.; Riaño, P.F.; Lugo, D.L.; Barriega, J.A.; Celis, C.A.; Moreno, S.; Pérez, A. Degradation of crystal violet by Catalytic Wet Peroxide Oxidation (CWPO) with mixed Mn/Cu oxides. *Catalysts* **2019**, *9*, 530. [[CrossRef](#)]
9. Lorenzo, D.; Dominguez, C.M.; Romero, A.; Santos, A. Wet peroxide oxidation of chlorobenzenes catalyzed by goethite and promoted by hydroxylamine. *Catalysts* **2019**, *9*, 553. [[CrossRef](#)]
10. Magioglou, E.; Frontistis, Z.; Vakros, J.; Manariotis, I.D.; Mantzavinos, D. Activation of persulfate by biochars from valorized olive stones for the degradation of sulfamethoxazole. *Catalysts* **2019**, *9*, 419. [[CrossRef](#)]
11. Messele, S.A.; Bengoa, C.; Stüber, F.E.; Giralt, J.; Fortuny, A.; Fabregat, A.; Font, J. Enhanced degradation of phenol by a fenton-like system (Fe/EDTA/H₂O₂) at circumneutral pH. *Catalysts* **2019**, *9*, 474. [[CrossRef](#)]



© 2019 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (<http://creativecommons.org/licenses/by/4.0/>).