

N°1100 / PC

TOPIC(s) : Biomass conversion / Homogenous, heterogenous and biocatalysis

Catalytic oxidation of lignin into aromatic compounds using CuO/TiO₂

AUTHORS

Antonio HERNANDEZ MANAS / IRCELYON, CNRS/UCBL-UNIVERSITÉ DE LYON, 2 AVENUE ALBERT EINSTEIN, VILLEURBANNE

Pascal FONGARLAND / CNRS/CPE LYON/UCBL-UNIVERSITE DE LYON, 43, BD DU 11 NOVEMBRE 1918 B.P. 82077, VILLEURBANNE

Léa VILCOCQ / CNRS/CPE LYON/UCBL-UNIVERSITE DE LYON, 43, BD DU 11 NOVEMBRE 1918 B.P. 82077, VILLEURBANNE

Laurent DJAKOVITCH / IRCELYON, CNRS/UCBL-UNIVERSITÉ DE LYON, 2 AVENUE ALBERT EINSTEIN, VILLEURBANNE

Corresponding author : Laurent DJAKOVITCH / laurent.djakovitch@ircelyon.univ-lyon1.fr

PURPOSE OF THE ABSTRACT

The current situation of resources depletion and increasing pollution has increased the research in the fields of biomass and lignin valorisation. Lignin is produced in large quantity in the paper industry that makes it an interesting alternative resource for producing new high added-value molecules. Several interesting compounds can be produced from lignin, mainly aromatics since it is the only biopolymer composed by aromatic monomeric units. In this work, a heterogeneous copper-based catalyst has been used to optimize lignin oxidative depolymerization to maximize aromatic yields.

Technical lignins obtained from different biomasses have been studied. They includes Softwood (Resinous and Pine), Hardwood (Eucalyptus) and Wheat straw. Lignins were extracted by several methods as Kraft, Organosolv and Soda processes. Catalytic oxidizing depolymerisation were conducted in alkaline conditions in the presence of heterogeneous copper-based catalyst. The reaction conditions were stated to 150°C, 20 Bar of air and 1800 rpm. 5%CuO/TiO₂ has been used as catalyst. Three main fractions have been obtained from this process: the residual lignin that exhibit various chemical modifications, an aromatic fraction mainly composed by vanillin, acetovanillone and syringaldehyde, and an aliphatic fraction mainly composed by succinic, oxalic and acetic acids.

The catalyst have been synthetized by incipient wetness impregnation, and calcined at 650°C. Vanillin is one of the main aromatic compounds produced. Using CuO/TiO₂ catalyst helped to increase the vanillin production from 3 to 5%. A maximal vanillin production was reached after 30 minutes of reaction time. Hardwood and Wheat straw lignins show higher degrees of oxidation producing an important amount of aliphatic fraction (68-55%) compared to softwood lignins (33-46%). Despite higher transformation, the aromatic fraction ratio was not increased. As expected from these biomasses, it showed an important quantity of syringaldehyde. Next, we observed that the extraction method played also an important role. As an example, for Wheat straw, Organosolv extraction led to increased aliphatic fraction ratio with reduced quantity of aromatics compared to Soda process. Similar observations comparing Kraft and Organosolv processes were made. The reaction conditions are being optimized to maximize the aromatic production. All the results will be presented and discussed in more detail.

Acknowledgement: The authors thanks Evonik for gift of TiO₂ support for catalyst preparation.

FIGURES

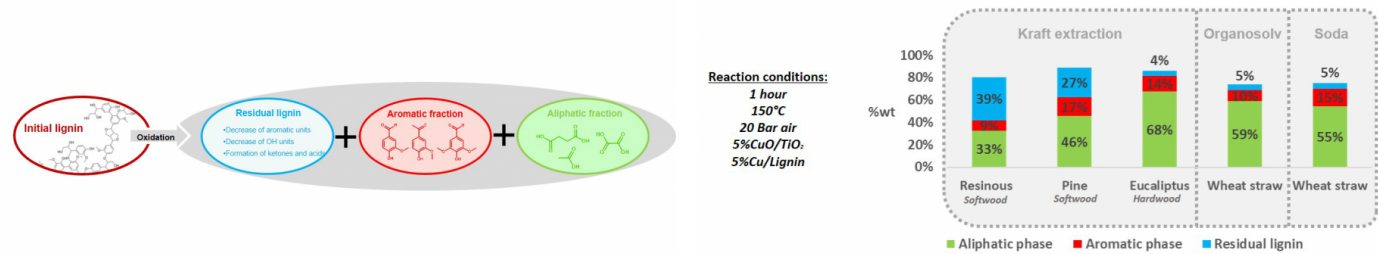


FIGURE 1
Concept of the Phenoliq project

FIGURE 2
Products fractionation after oxidative depolymerization of technical lignins
Catalyst CuO/TiO₂, 150°C, 60 min, 20 bar air.

KEYWORDS

Technical lignin | Oxidation reaction | Copper catalyst | biosourced phenols

BIBLIOGRAPHY

[1] F. G. Sales, L. C. A. Maranhão, N. M. L. Filho, and C. A. M. Abreu, "Experimental evaluation and continuous catalytic process for fine aldehyde production from lignin," Chem. Eng. Sci., vol. 62, no. 18–20, pp. 5386–5391, 2007.

[2] Villar, J. C., Caperos, A. (1997). Oxidation of Hardwood Kraft- Lignin to Phenolic Derivatives . Nitrobenzene and Copper Oxide as Oxidants. Journal of Wood Chemistry and Technology 17(3), 259–285.

[3] Ma, R., Guo, M. and Zhang, X. (2018). Recent advances in oxidative valorization of lignin. Catalysis Today. Elsevier 302, 50–60.