



Slovak Society of Chemical Engineering  
Institute of Chemical and Environmental Engineering  
Slovak University of Technology in Bratislava

## PROCEEDINGS

51<sup>st</sup> International Conference of the Slovak Society of Chemical Engineering SSCHE 2025

Hotel DRUŽBA  
Jasná, Demänovská Dolina, Slovakia  
May 27 - 30, 2025

Editors: Assoc. Prof. Mário Mihaľ

ISBN: 978-80-8208-158-2, EAN: 9788082081582

Published by the Faculty of Chemical and Food Technology Slovak Technical University in Bratislava in Slovak Chemistry Library for the Institute of Chemical and Environmental Engineering; Radlinského 9, 812 37 Bratislava, 2024

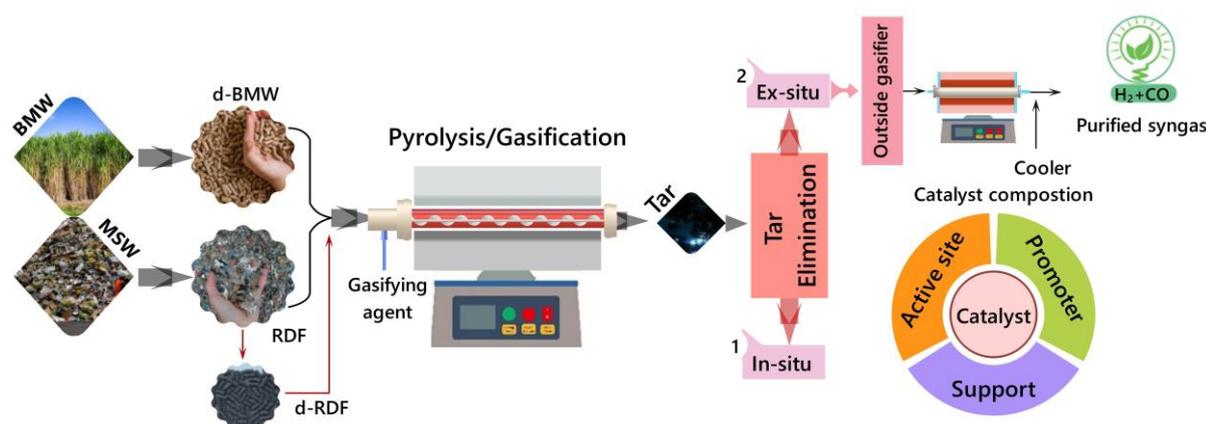
Nazari, M., Koritár, M., Haydary, J.: Ex-situ tar elimination during densified refuse-derived fuel (d-RDF) gasification, Editors: Mihaľ, M., In *51st International Conference of the Slovak Society of Chemical Engineering SSCHE 2025*, Jasná, Demänovská Dolina, Slovakia, 2025.

## Ex situ tar elimination during densified refuse-derived fuel (d-RDF) gasification.

Mohammad Ali Nazari, Matej Koritar, Juma Haydary

Institute of Chemical and Environmental Engineering, Slovak University of Technology in Bratislava, Radlinskeho 9, 812 37 Bratislava

### Graphical abstract



### Abstract

The formation of tar presents a major obstacle to the advancement of densified refuse-derived fuel (d-RDF) pyrolysis/gasification technologies, highlighting the efficient tar conversion strategies critically important. Catalytic tar reforming has emerged as a prominent technique for improving the efficiency of d-RDF gasification. Using activated carbon (AC), with abundant pore structure, high surface area, and unique chemical properties demonstrated an efficient potential as a support in tar conversion, enhancing resource application and system efficiency. However, catalyst deactivation is a widespread issue that remains a key challenge in catalytic tar conversion, significantly limiting its industrial application. This paper investigates AC modification, activation, catalytic performance in tar reforming, deactivation mechanisms, and regeneration strategies. The ex-situ catalytic performance of the catalysts was evaluated at various reaction temperatures using a two-stage fixed-bed reactor. The Ni/AC catalyst, with a specific surface area of 925.06 m<sup>2</sup>/g, demonstrated promising performance. As the catalyst loading increased from 0 to 18 g, the tar content was significantly reduced from 15 g/Nm<sup>3</sup> to 2.1 g/Nm<sup>3</sup>, achieving a high tar conversion efficiency of 85 % at a catalytic reaction temperature of 800 °C. H<sub>2</sub> was the dominant gas in the product gas, likely due to the enhanced cracking and reforming reactions of tar facilitated by the catalysts. The total gas yield reached 53.5 %, with a lower heating value (LHV) of 12 MJ/Nm<sup>3</sup>, H<sub>2</sub> yield of 13 mol/kg d-RDF, and a CO yield of 2.9 mol/kg d-RDF. In comparison to previous studies, the Ni/AC catalyst demonstrated competitive performance in terms of tar conversion efficiency. However, despite these advancements, sintering and coke deposition remain significant challenges in catalytic conversion of tar, necessitating improved modification strategies to enhance catalyst stability and activity.

**Keywords:** gasification, tar formation, catalyst, ex-situ tar conversion.