Catalytic performance and stability of Ru on Ce-based aminoclay carriers for Sabatier reaction

P1.49

A. Kaloudi¹, C. Drosou², K. Spyrou¹, P. Zygouri¹, I. V. Yentekakis^{2.3}, D. P. Gournis^{1*}

¹Department of Materials Science and Engineering, University of Ioannina, Ioannina, Greece.

²School of Chemical & Environmental Engineering, Technical University of Crete, Chania, Greece.

³Institute of GeoEnergy, Foundation for Research and Technology-Hellas (FORTH/IG), Chania, GR.

E-mail: a.kaloudi@uoi.gr, dgourni@uoi.gr

Keywords: CO2 methanation; Sabatier reaction; Ru; Ce aminoclay and La-Ce aminoclay

Carbon dioxide (CO₂) is a colorless, and odorless greenhouse gas (GHG) with a faint acid taste. It can be found naturally in springwater, etc., while it is the main anthropogenic pollutant from fossil fuel burn. Over the past years, CO₂ concentrations in atmosphere have increased by 30%. As a result, the global surface temperature increased from 0.4 to 0.8 °C.^[1]

The catalytic hydrogenation of CO₂ to produce CH₄, also known as CO₂ methanation or Sabatier reaction (CO₂ + 4H₂ ↔ CH₄ + 2H₂O; ΔHo= -164.7 kJ/mol) is considered an extremely important route for CO₂ recycling. The reaction is exothermic, thermodynamically favored at low temperatures (ca. 200- 400 °C), however, kinetically controlled (limited) at this region. Among various metals, Ni and Ru are among the most active Sabatier catalysts and have been extensively studied in an attempt to further promote their activity through metal–support interactions or other promotion strategies. [2.3]

Due to their unique combination of swelling behavior, intercalation, and ion exchange properties, Synthetic AminoClay Analogues (SACA or AC) have been employed in many application fields including catalysis, separation, gas adsorption, nanocomposites, ferrofluids, biomedicine etc.^[4–6] These tailor-made synthetic materials are layered magnesium organosilicates with structure analogous to 2:1 natural phyllosilicate smectite clays with covalently linked organosilicates in place of inorganic silicates, with an approximate composition R₈Si₈Mg₆O₁₆(OH)₄, where R is an organic functionality.^[7,8]

In this study, cerium, and lanthanum-cerium synthetic aminoclay analogues with an approximate composition R₈Si₈M₆O₁₆(OH)₄, where R represents propylamine functionalities (R = -CH₂CH₂CH₂NHCH₂CH₂NH₂), and M the cerium and lanthanum/cerium (1:1) metals were successfully prepared (CeAC and La-CeAC). The preparation process is a green, facile, and cost-effective room-temperature sol-gel-based synthetic method using metal chloride and aminotrialkoxysilane precursors. Using these cerium and lanthanum/cerium based synthetic Aminoclay analogues as carriers of Ru nanoparticles (Ru/CeAC, Ru/La-CeAC), the performance of this new class of catalysts

POSTER SESSION 1: Monday 18 September "Materials for energy & sustainability"

was investigated under CO₂ hydrogenation conditions and were found to be highly active and selective towards CH₄ production.

FTIR spectra ensure the successful synthesis of the CeAC and La-CeAC, while XRD patterns confirm the layered structure of the materials (Figure 1). As catalytic systems, Ru/CeAC and Ru/La-CeAC exhibit good CO₂ methanation and TOS performance. The up to 80% CH₄ yield obtained reveals that Ru/CeAC can be successfully used as efficient CO₂ methanation catalysts (Figure 2).

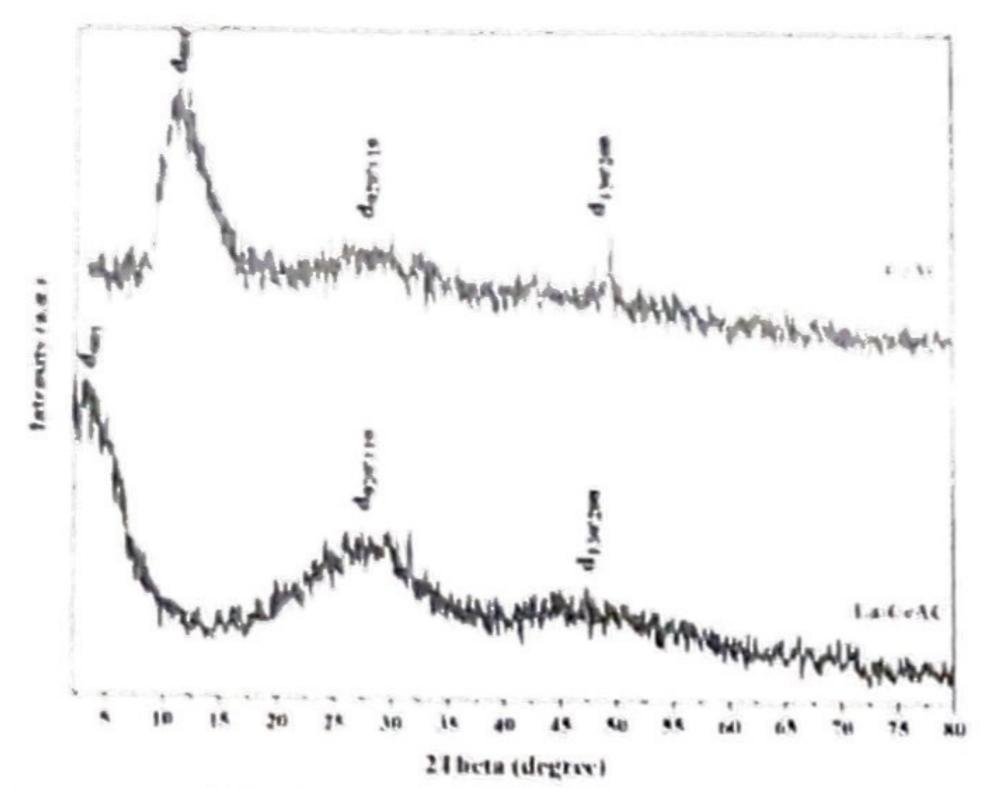


Figure 1. XRD patterns of CeAC and La-CeAC.

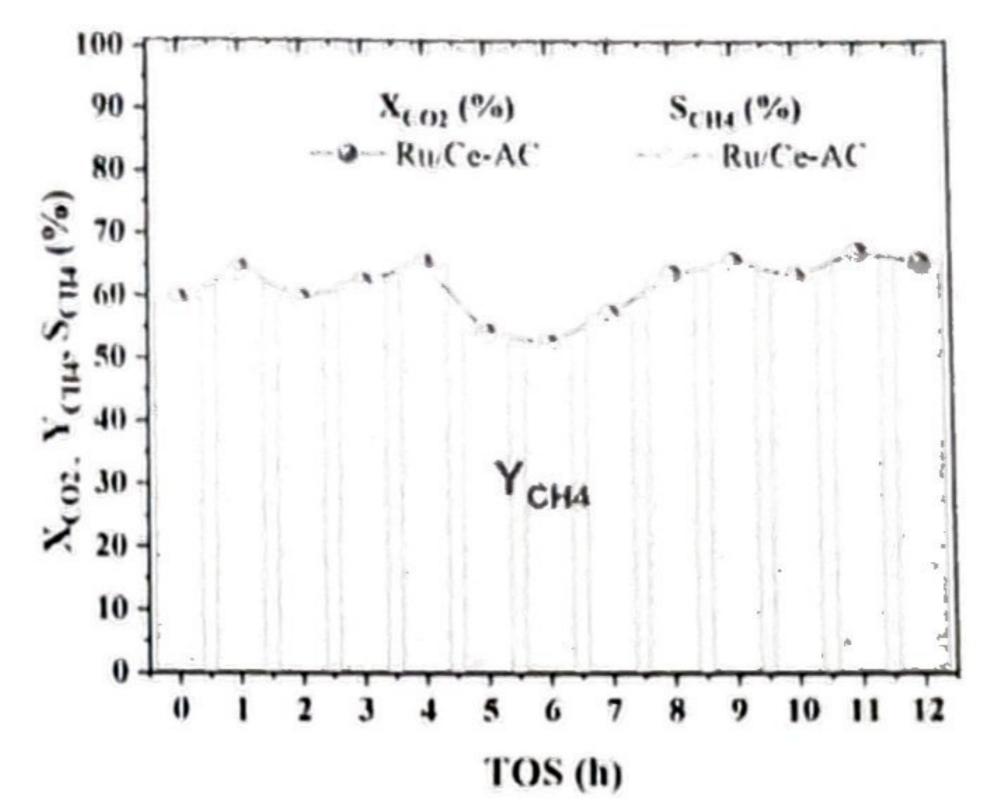


Figure 2. CO₂ conversion (X_{CO2}), CH₄ yield and selectivity (Y_{CH4} & S_{CH4}) for catalysts Ru/Ce-AC versus TOS (Time-on-stream, h). (T= 380°C).

Acknowledgements

The present work was co-funded by the European Union and Greek national funds through the Operational Program "Competitiveness, Entrepreneurship and Innovation", under the call "RESEARCH-CREATE-INNOVATE" (project code: Τ2ΕΔΚ-00955).

References

- C. H. Tan, S. Nomanbhay, A. H. Shamsuddin, Y.-K. Park, H. Hernández-Cocoletzi, P. L. Show, Front. Energy Res. 9 (2022) 795423.
- 2. J. Ashok, S. Pati, P. Hongmanorom, Z. Tianxi, C. Junmei, S. Kawi, Catal. Today 356 (2020) 471.
- 3. G. Botzolaki, G. Goula, A. Rontogianni, E. Nikolaraki, N. Chalmpes, P. Zygouri, M. Karakassides, D. Gournis, N. Charisiou, M. Goula, S. Papadopoulos, I. Yentekakis, Catalysts 10 (2020) 944.
- 4. V. K. H. Bui, D. Park, Y.-C. Lee, Chem. Eng. J. 336 (2018) 757.
- 5. L. P. Cavalcantí, G. N. Kalantzopoulos, J. Eckert, K. D. Knudsen, J. O. Fossum, Sci. Rep. 8 (2018) 11827.
- 6. L. Fu, K. K. R. Datta, K. Spyrou, G. Qi, A. Sardar, M. M. Khader, R. Zboril, E. P. Giannelis, Appl. Mater. Today 9 (2017) 451.
- 7. K. K. R. Datta, A. Achari, M. Eswaramoorthy, J. Mater. Chem. A 1 (2013) 6707.
- 8. K. O. Moura, H. O. Pastore, Environ. Sci. Technol. 47 (2013) 12201.