

CO₂ methanation: catalysts for CCU technologies

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Carbon dioxide capture, conversion, and storage (CCCS) technologies are emerging as pivotal solutions for utilizing carbon dioxide as a C1 building block¹. An intriguing solution lies in CO₂ methanation, leveraging existing methane grids globally and a clear regulatory framework. In our study, carbon dioxide methanation experiments were performed with two different setups, respectively operated at atmospheric and high-pressure. The former was used to conduct the catalyst screening while the latter was used to perform high-pressure tests and GHSV studies in a SS-316 tubular reactor. He was used as a diluent gas and N₂ was used as GC internal standard. The influence of Ni-loading, supporting materials and catalyst synthesis technique on the catalytic activity were investigated. Several oxides, such as ZrO₂, TiO₂, MgO, Al₂O₃, ZSM-5, SiO₂ and CeO₂, were tested as supports. CeO₂ showed extremely interesting performances, achieving high CO₂ conversion rates at high GHSV, as shown in Figure 1. Different preparation methods, such as sol-gel, co-precipitation and wet impregnation were investigated for the preparation of the Ni/CeO₂ catalyst. Several catalysts exhibited promising performance and despite the relatively high loadings, Ni catalysts emerged as viable alternatives to noble metal-based catalysts.

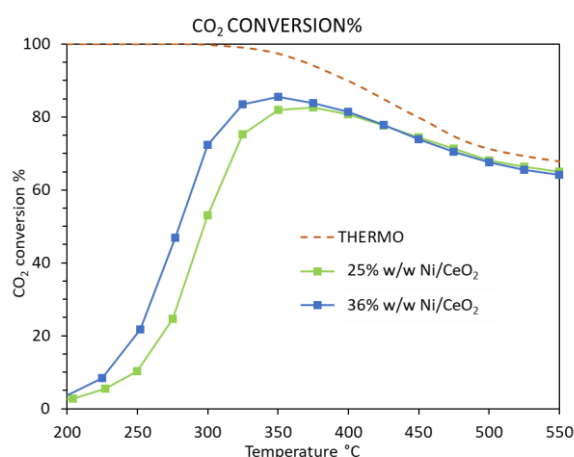


Figure 1. CO₂ conversion as function of the temperature for Ni/CeO₂ catalysts

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