




Dry reforming of methane for syngas production over Ni–Co-supported Al₂O₃–MgO catalysts

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Abstract

This research project focuses on the development of catalysts for syngas production by synthesizing Ni–Co bimetallic catalyst using aluminum oxide (Al₂O₃) and magnesium oxide (MgO) as the catalyst support. Ni/Al₂O₃ (CAT-1), Ni–Co/Al₂O₃ (CAT-2) and Ni–Co/Al₂O₃–MgO (CAT-3) nanocatalysts were synthesized by sol–gel method with citric acid as the gelling agent, and used in the dry reforming of methane (DRM). The objective of this study is to investigate the effects of Al₂O₃ and MgO addition on the catalytic properties and the reaction performance of synthesized catalysts in the DRM reactions. The characteristics of the catalyst are studied using field emission scanning electron microscope (FESEM), Brunauer–Emmett–Teller (BET), X-ray powder diffraction (XRD), transmission electron microscopy, H₂-temperature programmed reduction, CO₂-temperature programmed desorption and temperature programmed oxidation analysis. The characteristics of the catalyst are dependent on the type of support, which influences the catalytic performances. FESEM analysis showed that CAT-3 has irregular shape morphology, and is well dispersed onto the catalyst support. BET results demonstrate high surface area of the synthesized catalyst due to high calcination temperature during catalysts preparation. Moreover, the formation of MgAl₂O₄ spinel-type solution in CAT-3 is proved by XRD analysis due to the interaction between alumina lattice and magnesium metal which has high resistance to coke formation, leading to stronger metal surface interaction within the catalyst. The CO₂ methane dry reforming is executed in the tubular furnace reactor at 1073.15 K, 1 atm and CH₄/CO₂ ratio of unity to investigate the effect of the mentioned catalysts. Ni–Co/Al₂O₃–MgO gave the highest catalyst performance compared to the other synthesized catalysts owing to the strong metal–support interaction, high stability and significant resistance to carbon deposition during the DRM reaction.

Keywords Catalyst development · Dry reforming · Bimetallic catalysts · Catalysis · Sol–gel · Support

Introduction

Recently, the global warming issue is getting crucial due to the substantial dependence on petroleum-based energy that leads to the increment of greenhouse-gas emissions

within the atmosphere [30]. The concentration of CO₂ in the atmosphere has currently increased by about 1.5 ppm/year which indicated that if there is about 5.3×10^{21} g air, the rate of CO₂ increase is about 8 billion tons per year [5]. These realities encouraged the study on the development

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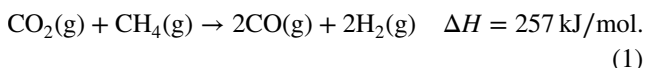
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of CO₂ reforming of methane that would effectively reduce the level of CO₂ and CH₄ within the atmosphere.

Steam reforming of methane (SRM), partial oxidation of methane (POM) and methane dry reforming (DRM) [7] are three common techniques which have been used in industries. SRM is used conventionally for hydrogen generation [17, 35]. However, a foreseeable drawback of this process is the production of H₂/CO with the ratio of 3:1 that is undesirable for Fischer–Tropsch (F–T) synthesis. CO₂ reformation of methane is based on the utilization of CO₂ and CH₄ to convert these gases into syngas with low or adjustable hydrogen–carbon monoxide ratio (H₂/CO), which is a desirable feedstock for F–T synthesis to gain liquid hydrocarbons [9, 10, 13]. Therefore, DRM has been considered as the favorable process for syngas production compared to steam reforming and partial oxidation, since it produces lower H₂/CO ratio that is appropriate for downstream F–T synthesis [13]. Yet, the industrial applications of DRM are still facing vast challenges in terms of process engineering and catalysts stability [20]. DRM involves high operating temperatures, usually between 900 and 1273 K to enhance the conversions, and low operating pressures to favor the forward reaction in DRM [6]. The DRM reaction is shown in Eq. (1):



Ni-based catalysts are the best alternatives for the noble metals to be used in the DRM reaction as they have high catalytic performances, wide availability and low cost [11, 24]. Eventhough Ni/Al₂O₃ catalyst has been utilized broadly as DRM reaction catalyst, it encounters a rapid catalyst deactivation that restricts its industrial application. Coke formation [9, 26, 28] and sintering of catalyst at high temperature [17, 35] are the main reasons of the catalyst deactivation. The coke formation primarily originates from two side reactions in DRM which are methane cracking reaction (Eq. 2) and Boudouard reaction (Eq. 3). Also, the highly endothermic nature of DRM reaction requires high temperatures to achieve the desirable conversions [6]. Thus, most of the catalysts are not thermally stable at these conditions due to the sintering of catalyst or collapse of the crystal structure [22]. Hence, it is important to develop a novel catalyst which meets these significant criteria such as high thermal stability of the support, better resistance to coke formation, and sintering of the active phase and high activity throughout the reaction time to achieve the equilibrium conversions [22]:



The development of cost-effective catalysts having higher catalytic activity and more considerable resistance

to carbon formation is one of the most concerning issues to commercialize DRM reaction in the industries [3]. One of the approaches applied for the DRM reaction to develop high carbon resistance of Ni-based catalyst is the addition of second metal [8]. The addition of a non-noble metal such as cobalt is more preferable from the economic point of view [8]. Xu et al. [32] proved that the ratio of Ni/Co is closely allied to the catalytic activity of the bimetallic Ni/Co catalysts supported with the γ -Al₂O₃ and doped with La₂O₃. The catalyst having the Ni/Co ratio of 7/3 shows the greatest CH₄ and CO₂ conversion. If the amount of cobalt increases, the catalytic activity decreases. Bimetallic catalyst having ratio of 5/5 and 3/7 exhibits lower catalytic activity than monometallic Ni catalyst [32].

Conventionally, Al₂O₃ is the most proper support for most of the catalytic materials owing to mechanical strength, stability at high temperature and also good textural properties [31]. Although it has been commercially used, the coke deposition is one of the drawbacks of using Al₂O₃ due to its acidic properties. MgO and CeO₂, which are known as the alkaline and alkaline earth oxides, are being used as modifiers of Ni-based catalysts [18] to enhance the metallic dispersion, improve the metal–support interaction, reduce sintering and improve the thermal stability [23, 27]. The basicity of the catalyst is predicted to be increased by the incorporation of MgO and CeO₂. The catalyst basicity improves the adsorption of CO₂ which prevents the formation of coke on the catalyst surface [14, 29, 33].

In present work, the influence of bimetallic Ni–Co catalysts supported with Al₂O₃ and Al₂O₃–MgO in DRM reaction, synthesized by the sol–gel method, is being studied. The synthesized catalysts were characterized and analyzed in tubular furnace reactor to explain the effect of supports on reaction performances to enhance the reactant conversions as well as to minimize the coke formation in DRM reaction.

Experimental section

Materials

The materials used for the catalysts preparation were Ni(NO₃)₂·6H₂O (EMSURE ACS, 99%) and Co(NO₃)₂·6H₂O (HmbG Chemicals, 97%) as the active metals; Al(NO₃)₃·9H₂O (HmbG Chemicals, 98.5%) and Mg(NO₃)₂·6H₂O (EMSURE ACS, 99%) as the catalyst supports; and citric acid as the gelling agent. High-purity CO₂, CH₄, N₂ and H₂ (Linde) were used as the laboratory gases for the reaction.

Catalysts preparation

The new Ni–Co bimetallic catalysts were synthesized by direct sol–gel method [19] as it has high potential in producing high homogeneity composition, and improves the particle size distribution in nanoscale levels that could lead to the high catalytic performance [1, 16]. The $\text{Ni}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$, $\text{Co}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$, $\text{Al}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$, $\text{Mg}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$, and citric acid were dissolved in deionized water. The continuously stirred mixture was heated using hot plate at 333.15 K until the gel was formed. Consequently, the resulting gel was dried in an oven at 383.15 K overnight, and then calcined in the furnace at 1173.15 K for 5 h. The other samples were also been prepared by the above procedure.

Table 1 includes the formation of the catalysts, including their tagging, molecular structure, description and composition.

Catalyst characterization

FESEM was employed to determine the morphological change by scanning the catalyst samples with a high-energy beam of electrons using a Zeiss Supra_55 VP. Brunauer–Emmett–Teller (BET) analysis was used to determine the specific surface area of a sample—including the pore size distribution. 0.5 g catalyst was used for each analysis using a Micromeritics ASAP 2020. The degassing temperature was set at 383 K to remove the moisture and other adsorbed gases from the catalyst surface.

Furthermore, the phase compositions of the synthesized catalysts were defined by X-ray powder diffraction (XRD) analysis using a Bruker D8B advanced X-ray diffractometer which were recorded in the range $2\theta = 20^\circ - 80^\circ$. The crystallite size, t , was estimated from X-ray line broadening using the Scherrer's formula, $t = 0.9 \cdot \lambda / (B \cos(\theta))$, where λ is the X-ray wavelength (Cu $K\alpha$ radiation 0.154 nm) and B the full-width half-maximum of the Bragg diffraction angle θ . To study the metal dispersion on the catalyst support, transmission electron microscopy (TEM) was conducted using a HITACHI instrument which operated at 120.0 kV.

Moreover, the reducibility performance of the synthesized catalysts was determined by H_2 -temperature programmed reduction (H_2 -TPR) analysis technique on a Thermo Finnigan (TPDRO 1100) instrument equipped with a thermal conductivity detector (TCD) in two-stage processes, which

are pretreatment and analysis. CO_2 -temperature programmed desorption (CO_2 -TPD) was used to study the basic properties of the synthesized catalysts. The result is gained from the TP-5000 equipment coupled with a Hiden QIC-20 mass spectrometer. For post-reaction analysis, TPO was used with the same equipment and procedures of H_2 -TPR. 0.2 g of the spent catalysts and 5% of O_2/He gas mixture were introduced for the TPO analysis.

Catalytic performance evaluation and testing

The performance of Ni-based supported catalysts in the DRM was studied in a tubular furnace reactor under atmospheric pressure. 0.2 g of the catalyst was held in the middle of the reactor tube between two layers of quartz wool, and the reactor was electrically heated in a furnace. The reactor was purged with N_2 gas at 100 mL min^{-1} to provide an inert atmosphere in the reactor prior to starting the experiment. The reduction process was then carried out at a H_2 flow rate of 20 mL min^{-1} and at a temperature of 1023.15 K for 1 h to activate the catalyst. Then, nitrogen gas was purged again in the reactor until the gas chromatography system showed a complete disappearance of hydrogen gas before the reaction initiation. Methane gas and carbon dioxide with a flow rate of 20 mL min^{-1} for each gas were introduced to the reactor for every run. The reaction took 8 h for every run, and the sample was taken every hour. The effluent from the reactor then was analyzed by an online gas chromatograph (Agilent 7890) with a thermal conductivity reactor (TCD) and a flame ionization detector (FID).

Results and discussion

Morphology

Figure 1 shows the FESEM images of the synthesized catalysts after calcination. CAT-1 and CAT-2 reveal similar morphologies which are the long-shaped nanoparticles. CAT-3 which is Ni–Co/ Al_2O_3 –MgO shows a clear irregular-shaped morphology image which has a better dispersion onto the catalyst support compared to CAT-2 with a broad size distribution ranging from 20 to 80 nm. The addition of MgO support improves the metal particles dispersion. This leads

Table 1 Mass composition for synthesized catalysts

Catalyst tagging	Catalyst	Description of catalyst	Composition of catalyst
CAT-1	Ni/ Al_2O_3	Support: Al_2O_3	15% Ni, 85% Al_2O_3
CAT-2	Ni–Co/ Al_2O_3	Support: Al_2O_3	10.5% Ni, 4.5% Co, 85% Al_2O_3
CAT-3	Ni–Co/ Al_2O_3 –MgO	Support: Al_2O_3 and MgO	10.5% Ni, 4.5% Co, 63.75% Al_2O_3 , 21.25% MgO

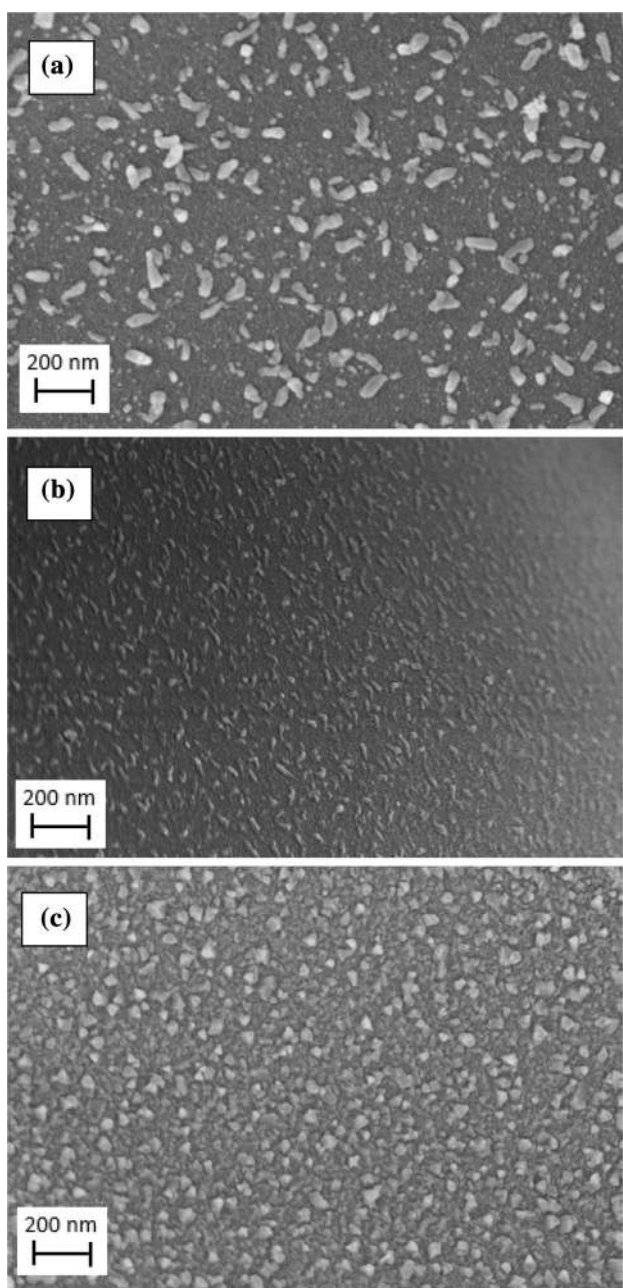


Fig. 1 FESEM images of catalysts at 30.0Kx magnification. **a** CAT-1; **b** CAT-2; **c** CAT-3

to more active sites available for the reaction, and enhances the catalytic performance.

TEM analysis

Figure 2 indicates the TEM analysis results for the synthesized catalysts. Due to the large particles formation, CAT-1 and CAT-2 show a clear catalyst agglomeration. Moreover,

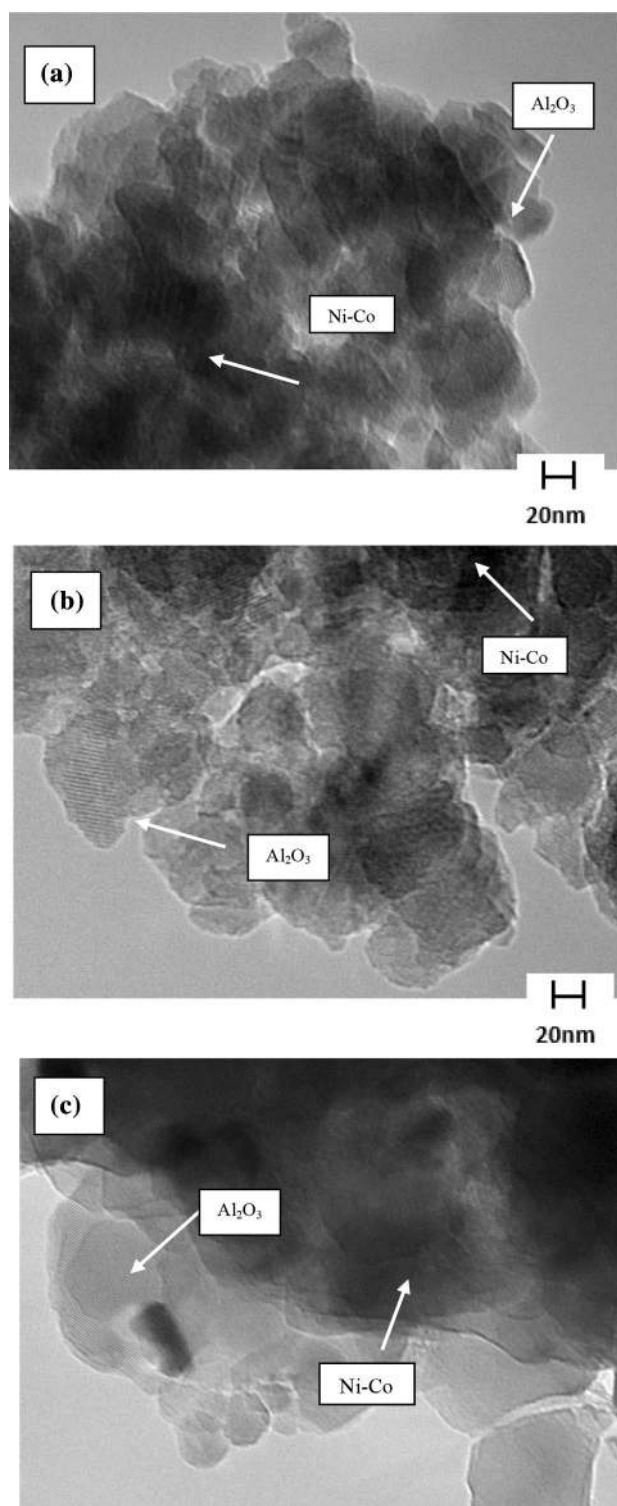


Fig. 2 TEM micrographs of the synthesized catalysts. **a** CAT-1; **b** CAT-2; **c** CAT-3

the TEM micrographs for CAT-3 demonstrates that there are more uniform NiO particles distribution on the catalyst surface that supports the previous FESEM result (cf. Figure 1).

Textural properties

The textural properties of the synthesized catalysts are summarized in Table 2. The surface area and the pore volume for CAT-1, which is Ni–Co/Al₂O₃, were found to be 21.58 m² g⁻¹ and 0.04 cm³ g⁻¹, respectively. Moreover, the surface area and pore volume for CAT-3, which is Ni–Co/Al₂O₃–MgO, were measured to be 9.76 m² g⁻¹ and 0.02 cm³ g⁻¹, respectively. Al-Fatash and Fakeeha [4] studied on the effect of calcination temperatures on DRM catalysts using Ni/Al₂O₃ and claimed that the high calcination temperature reduces the surface areas but has the advantage of getting the existence of a stable structure of the catalysts. This claim is supported with the BET results in which the catalysts have been calcined at high temperature (1173.15 K).

XRD analysis

Figure 3 illustrates the XRD patterns of the synthesized catalysts at 2θ = 20°–80°. The XRD results indicate five crystalline phases formation of MgAl₂O₄, Co₃O₄, NiO, MgO and γ-Al₂O₃. The MgAl₂O₄ peaks in the form of cubic phase are recognized at 31.6°, 37.3°, 45.3°, 60.0° and 66.2° (JCPDS 00-001-1157) for CAT-3, while Co₃O₄ diffraction peaks are identified at 2θ = 31.3°, 36.9°, 44.9°, 59.5°, and 65.3° (JCPDS 01-076-1802) for CAT-2 and CAT-3. The peaks for NiO from MgAl₂O₄ and MgO peaks, and peaks for Co₃O₄ from MgAl₂O₄ peaks are difficult to differentiate due to the existing overlaps. The figure signifies the patterns existence of NiO peaks at 37.3°, 43.4°, 63.0°, 75.6°, and 79.6° (JCPDS 01-073-1519). For MgO, the diffraction peaks are revealed at 2θ = 37.0°, 43.0°, 62.4°, 74.8°, and 78.7° (JCPDS 01-077-2364). The MgO diffraction peaks indicate the high degree of crystallinity in Al₂O₃–MgO support for CAT-3. Furthermore, γ-Al₂O₃ peaks are identified in Ni/Al₂O₃ catalyst at 2θ = 42.8° and 67.3° (JCPDS 00-004-0880) in cubic phase. This analysis indicates that there is an interaction between alumina lattice and magnesium metal to form MgAl₂O₄ spinel-type solid solution. MgAl₂O₄ has high resistance to carbon formation, and can enhance the metal surface interaction strength [2]. The crystallite sizes calculated using Scherrer's equation are 18.34 nm, 13.1 nm, and 10.73 nm for CAT-1, CAT-2 and CAT-3, respectively. From these observations,

Table 2 Textural properties for synthesized catalysts

Catalyst	BET _s (m ² /g)	Pore volume (cm ³ /g)	Pore size (Å)
CAT-1	25.57	0.03	48.32
CAT-2	21.58	0.04	57.81
CAT-3	9.76	0.02	107.87

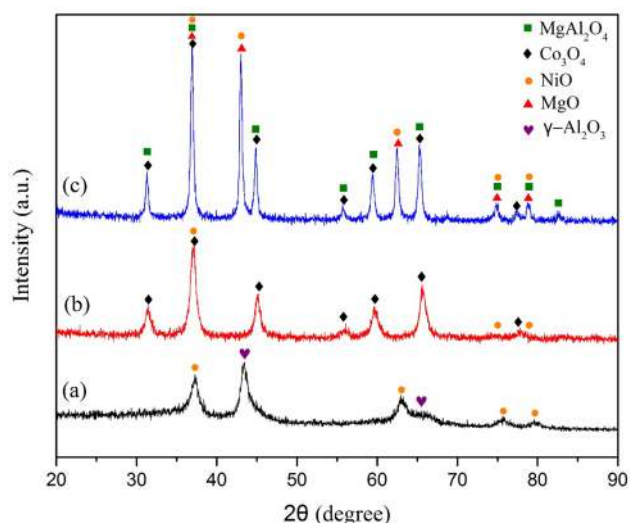


Fig. 3 X-ray diffraction patterns of the catalysts prepared by sol-gel method. (a) CAT-1; (b) CAT-2; (c) CAT-3

it can be concluded that CAT-3 has a better dispersion compared to the other synthesized catalysts, which is supported by the FESEM results (cf. Fig. 1).

H₂-TPR analysis

Figure 4 displays the H₂-TPR analysis for all synthesized catalysts. The TPR patterns are observed to be similar for all synthesized catalysts which have only one peak. The peaks for CAT-1, CAT-2, and CAT-3 are observed at 1180.15 K, 1040.15 K, and 1170.15 K, respectively. These peaks are formed owing to the reduction of complex NiOx species

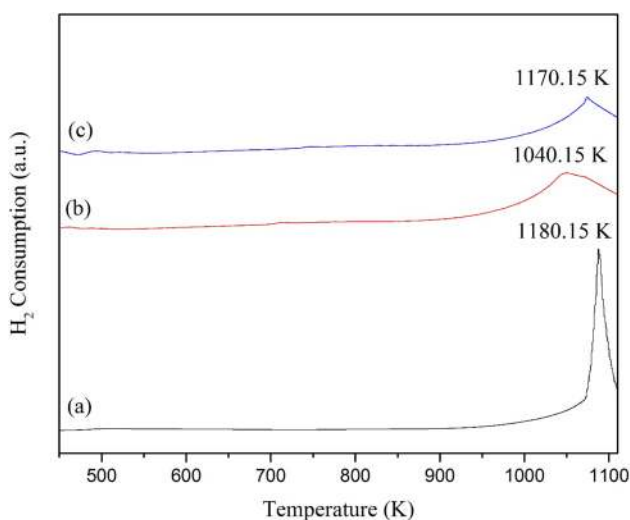


Fig. 4 H₂-TPR patterns of the synthesized catalysts. (a) CAT-1; (b) CAT-2; (c) CAT-3

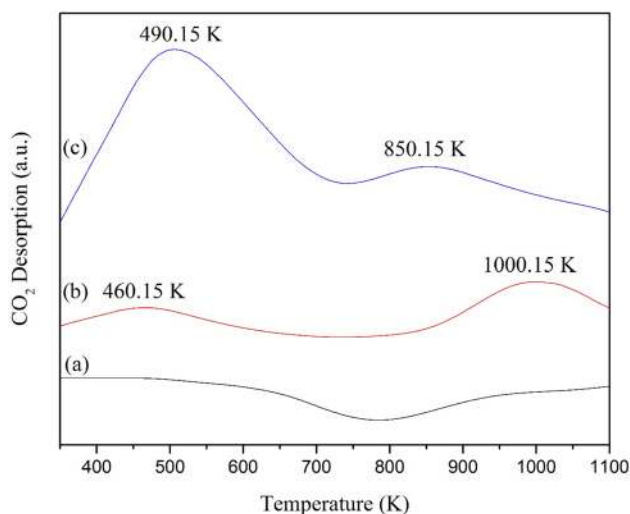


Fig. 5 CO₂-TPD patterns of the synthesized catalysts. (a) CAT-1; (b) CAT-2; (c) CAT-3

which has a strong interaction between the active metals and catalyst support [21].

CO₂-TPD analysis

The CO₂ desorption patterns shown in Fig. 5 display the weak base sites' peak around 490.15 K and the strong base sites' peaks around 850.15 K. In CAT-3, the desorption temperature shifted to the higher location due to the formation of MgAl₂O₄ that has strong basic properties. The amount of CO₂ absorbed is increased, and offers more oxygen species on the surface of the catalyst which is useful for the DRM reaction [12]. Hence, an increase in the supports' Lewis basicity could result in a coke-free DRM reaction which boosts the capability of catalysts to chemisorb CO₂, which reacts with C to form CO [12]. Also, this result supports the discussion in XRD analysis implying that the MgAl₂O₄ formation poses high resistance to carbon formation and strong metal–surface interaction which could improve the conversion of reactants.

Catalytic performance in DRM

An ideal catalyst for DRM reaction should have two significant properties which are the stability and high activity performance. Figure 6 illustrates the catalytic performances of CAT-1, CAT-2 and CAT-3 in the DRM reaction at 8-h time on stream. Moreover, there is an insignificant occurrence of the reverse water–gas shift (RWGS) reaction on the catalyst which is indicated by the H₂/CO ratio trend that is approximately close to 1, and it is in line with the results found by Zhang et al. [34].

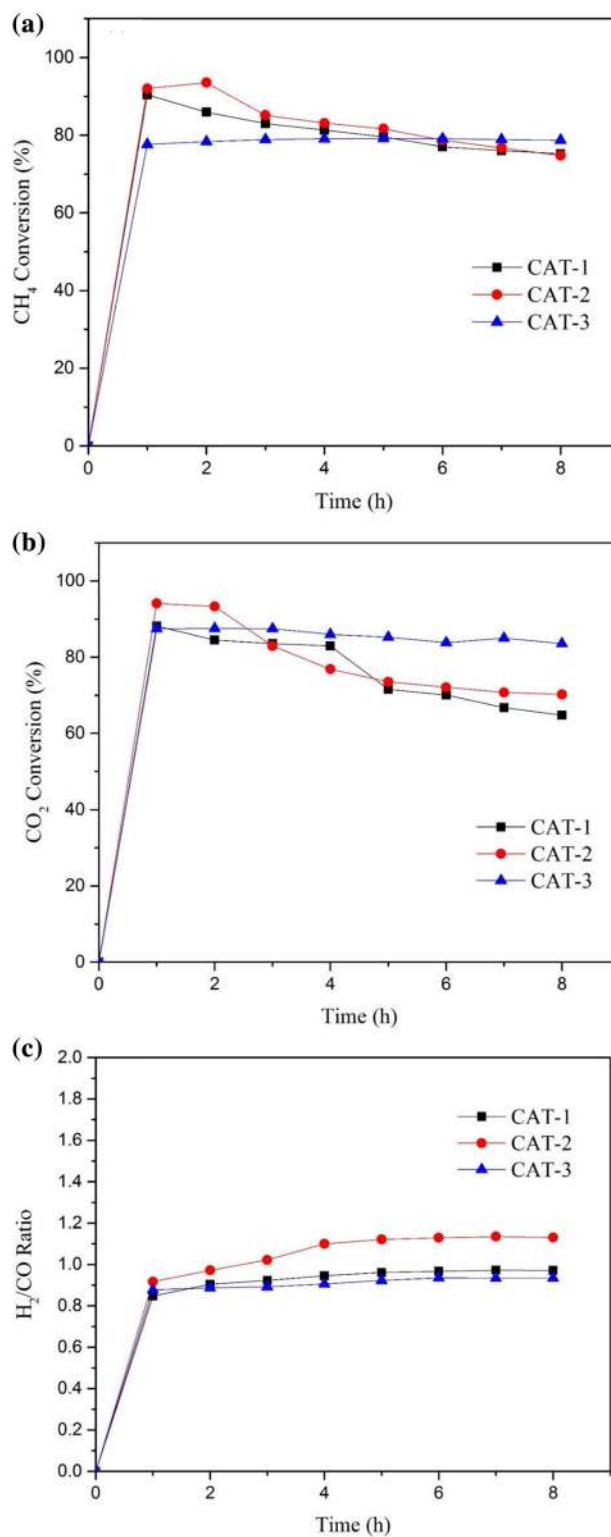


Fig. 6 The catalytic performance of **a** CH₄ conversion; **b** CO₂ conversion; and **c** H₂/CO ratio in DRM presented by synthesized catalysts under the following reaction conditions: $T=1073.15$ K, $P=1$ atm, $\text{CH}_4=20$ mL min⁻¹, $\text{CO}_2=20$ mL min⁻¹. Symbols represent: (black filled square line) CAT-1, (red filled circle line) CAT-2 and (blue filled triangle line) CAT-3

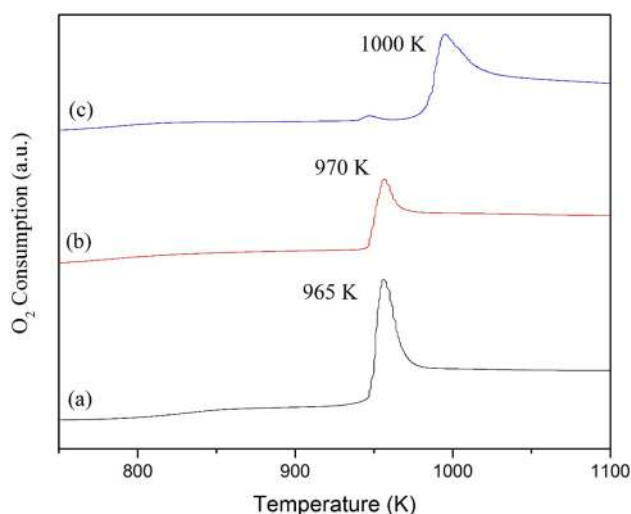


Fig. 7 Temperature programmed oxidation with 5% O₂ in He, performed after 8 h reaction

At the early reaction time, CAT-1 and CAT-2 have higher conversions for both CH₄ and CO₂ compared to CAT-3. However, the conversion of CAT-1 and CAT-2 keep on decreasing, implying that they are unstable throughout the reaction time. On the other hand, CAT-3 is the most stable but less active. This may be due to the low BET surface area of CAT-3 compared to the other two catalysts. However, the basic properties of MgO improve the support interaction and have high stability performance throughout the reaction. Also, the cooperation between Ni and Co results in high performance of catalysts in the DRM reaction. The employment of MgO also produces MgAl₂O₄ spinel type that enhances the carbon resistance of the catalyst. The average conversions of CH₄ and CO₂ for CAT-3 are higher, which are 79.17% and 84.82%, respectively, compared to the conversion of CH₄ and CO₂, which were only 55.7% and 60.9%, obtained from the results of Ni/Al₂O₃ catalyst prepared by Min et al. [25] with the reaction temperature of 1073.15 K and pressure of 1 atm.

Post-reaction analysis

TPO was executed to investigate the coke formation in the DRM reaction for the duration of 8 h. The amount of carbon dioxide formed during TPO quantifies the amount of coke formed during the reaction, and the temperature needed to burn off the carbon is an indicator for the carbon bond strength with the catalyst surface. Figure 7 conveys the CO₂ production during TPO in 5% of O₂ in N₂ after DRM reaction. CAT-1 has the highest amount of coke formed, and a temperature of 1000 K is needed to oxidize most of it. The CO₂ evolution at greater than 723 K indicates the oxidation of whisker-type carbon that does not deactivate the nickel

surface but slightly causes a breakdown of the catalyst by pore plugging [15].

Conclusion

A comparison between the three synthesized Ni–Co bimetallic catalysts is accomplished. The addition of MgO as the catalyst support into the bimetallic catalyst supported Al₂O₃ decreases the BET surface area and pore volume as MgO has low surface area which causes pore-filling during the catalyst preparation. XRD analysis proves that there is a formation of MgAl₂O₃ spinel-type solid solution which has high resistance to coke formation and high metal–support interaction. Hence, CAT-3 that is Ni–Co/Al₂O₃–MgO gives the highest catalyst performance compared to the other synthesized catalysts due to the addition of MgO which enhances the metal–support interaction, and suppresses the carbon formation in DRM reaction that can lead to the high stability and activity performance of the catalysts.

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