Vacuum Thermal Treated Ni-CeO$_2$/SBA-15 Catalyst for CO$_2$ Methanation

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Abstract: Ni-CeO$_2$/SBA-15-V catalyst was prepared by the impregnation method with vacuum thermal treatment and used for CO$_2$ methanation reaction. Compared with Ni-CeO$_2$/SBA-15-air catalyst with thermal treatment in air, the reduced Ni-CeO$_2$/SBA-15-V catalyst with vacuum thermal treatment exhibited higher Ni dispersion and smaller Ni particle size. In CO$_2$ methanation reaction, the Ni-CeO$_2$/SBA-15-V catalyst was more active and selective than the Ni-CeO$_2$/SBA-15-air catalyst. The good activity and selectivity of Ni-CeO$_2$/SBA-15-V catalyst should be due to highly dispersed Ni in contact with small CeO$_2$ particles.

Keywords: nickel; CeO$_2$; SBA-15; vacuum thermal treatment; CO$_2$ methanation

1. Introduction

With the increasing consumption of fossil fuel, CO$_2$ released into the atmosphere has resulted in climate change and global warming, and the utilization of CO$_2$ has received much attention. CO$_2$ hydrogenation is an effective way to convert CO$_2$ into fuels and chemicals [1,2]. CO$_2$ methanation can convert CO$_2$ and renewable H$_2$ into methane, which is a promising process for CO$_2$ conversion and the storage of renewable H$_2$ [3].

CO$_2$ methanation is an exothermic reaction, and the low reaction temperature is favorable for the high equilibrium conversion of CO$_2$. Therefore, various catalysts have been developed for CO$_2$ methanation at low temperature. Noble metal-based catalysts, such as Rh [4], Ru [5], and Pd [6], and non-noble Co [7,8] and Ni-based catalysts [9,10] are the most studied CO$_2$ methanation catalysts. Although noble metal-based catalysts possess good activity for low-temperature CO$_2$ methanation, Ni-based catalysts are preferred in industrial processes due to the low cost, and they have been applied for CO methanation [11,12] and CO$_2$ methanation [13]. Ni supported on various metal oxides, such as SiO$_2$ [13], Al$_2$O$_3$ [14], CeO$_2$ [15], ZrO$_2$ [16], and TiO$_2$ [17], has been reported for CO$_2$ methanation. Ni supported on SBA-15, which is a kind of mesoporous SiO$_2$ with high specific surface area, has been reported for CO$_2$ methanation [18,19]. Adding CeO$_2$ to a Ni-based catalyst can enhance Ni dispersion and CO$_2$ adsorption and dissociation [20]. CeO$_2$-promoted Ni/SBA-15 catalyst has been used for CO$_2$ methanation reaction and exhibits excellent catalytic performance [10,19].

However, it is difficult to obtain highly dispersed Ni particles on SBA-15 support by the conventional impregnation method, and Ni particles are mainly located on the SBA-15 outer surface [19]. Various methods, such as two solvents [21], surfactant assistant [22], and ammonia
evaporation [23] methods have been used to improve the dispersion of Ni on SBA-15 support. Recently, vacuum thermal treatment was applied to synthesize highly dispersed Cu [24], Pt [25], and Pd [26] particles on mesoporous SBA-15. In the present work, the Ni-CeO$_2$/SBA-15 catalyst was prepared by the impregnation method with vacuum thermal treatment and used for CO$_2$ methanation. The vacuum thermal treated catalyst exhibited high dispersion of Ni and CeO$_2$ and showed high activity and selectivity for CO$_2$ methanation reaction.

2. Materials and Methods

2.1. Synthesis of Catalysts

SBA-15 was prepared by a hydrothermal method according to the literature [27]. First, 16.4 g of P123 (EO$_{20}$PO$_{70}$EO$_{20}$, Mw = 5800; Sigma Aldrich, St. Louis, MO, USA) was dissolved in 500 mL of a 2 M HCl aqueous solution. After stirring at 40 °C for 1 h, 34.0 g tetraethoxysilane, (TEOS, Sinopharm, Shanghai, China) was dropped into the solution. Then it was stirred at 40 °C for 4 h, and aged at 100 °C for 48 h in sealed Teflon bottles. SBA-15 was obtained after filtration, drying at 80 °C overnight, and calcination at 600 °C for 4 h (heating rate of 1 °C/min).

Then, 1 g of SBA-15 was impregnated with controlled amounts of Ni(NO$_3$)$_2$·6H$_2$O and Ce(NO$_3$)$_3$·6H$_2$O using 5 mL of ethanol as solvent and kept at room temperature for 24 h, then it was calcinated in air or vacuum at 600 °C for 4 h with a heating rate of 1 °C/min. The resulting samples calcinated in air and vacuum are referred to as Ni-CeO$_2$/SBA-15-air and Ni-CeO$_2$/SBA-15-V, respectively.

For comparison, using γ-Al$_2$O$_3$ (241 m$^2$/g) as support, a Ni-CeO$_2$/Al$_2$O$_3$ catalyst was prepared by impregnation method and calcinated in air as described above.

In these catalysts, the theoretical Ni and CeO$_2$ content was 10 wt % for both.

2.2. Characterization of Catalysts

N$_2$ isotherms were measured using an Autosorb-iQ analyzer (Quantachrome Instruments, Boynton Beach, FL, USA) at −196 °C, and the pore size distribution was calculated from the desorption branch by Barrett–Joyner–Halenda (BJH) method. X-ray diffraction (XRD) patterns were measured by a DX-2700 X-ray diffractometer (Haoyuan Instrument, Dandong, China) using Cu K$_\alpha$ radiation. Transmission electron microscope (TEM) images were measured on a Tecnai G2 F20 microscope (FEI Company, Hillsboro, OR, USA). (XRD). H$_2$-temperature programmed reduction (H$_2$-TPR) patterns were measured on a TP-5080 adsorption instrument under a 5% H$_2$/Ar flow at a rate of 10 °C/min to 900 °C. Before H$_2$-TPR measurements, the samples were purged with an argon flow at 400 °C for 5 min. The outgas was monitored by a thermal conductivity detector (TCD) detector. Thermo gravimetric (TG) was carried out using an HCT-1 TG thermal analyzer (Henven Scientific Instrument, Beijing, China). The sample was heated from room temperature to 900 °C with a heating rate of 10 °C/min in air (20 mL/min).

2.3. Catalytic Test

The reaction was carried out in a fixed bed quartz reactor (inner diameter (ID) = 8 mm) at atmospheric pressure. The temperature of the reactor was controlled by a thermocouple inserted into the catalyst bed, and the gas flow rates were controlled by mass flow controllers. For a catalytic test, 100 mg of catalyst (60–100 mesh) diluted with 200 mg of inert SiO$_2$ was used. The Ni-CeO$_2$/SBA-15 catalyst was reduced to 50 mL/min of 20% H$_2$/Ar stream at 450 °C for 40 min, and. After reduction, the reactor temperature was decreased to 200 °C in 20% H$_2$/N$_2$. Then the reactant gases (CO$_2$:H$_2$:Ar = 1:4:5, 100 mL/min) were introduced into the reactor. The reaction was conducted in the temperature range from 200 to 450 °C. The catalyst was kept for 1 h at each reaction temperature before the products were analyzed. The stability tests were performed at 350 and 400 °C. Before stability test, the Ni-CeO$_2$/Al$_2$O$_3$ catalyst was reduced at 700 °C for 40 min. The exhaust and feed gas compositions
were analyzed on a GC-7900 gas chromatograph (Techcomp, Kwai Chung, China) equipped with a TDX-01 column and a TCD detector. CO₂ conversion, CH₄ selectivity, and carbon balance were calculated using the following formulae:

\[
\text{CO}_2 \text{ conversion} = \frac{\text{moles of CO}_2 \text{ in} - \text{moles of CO}_2 \text{ out}}{\text{moles of CO}_2 \text{ in}} \times 100\%
\]

\[
\text{CH}_4 \text{ selectivity} = \frac{\text{moles of CH}_4 \text{ out}}{\text{moles of CO}_2 \text{ in} - \text{moles of CO}_2 \text{ out}} \times 100\%
\]

\[
\text{Carbon balance} = (1 - \frac{\text{moles of CH}_4 \text{ out} + \text{moles of CO}_2 \text{ out}}{\text{moles of CO}_2 \text{ in}}) \times 100\%
\]

In this work, carbon balance was within ±3% for all the catalytic tests, indicating negligible carbon deposition on the catalysts.

3. Results and Discussion

3.1. Characterization of Catalysts

Figure 1 shows N₂ adsorption–desorption isotherms and pore size distributions of SBA-15, Ni-CeO₂/SBA-15-air, and Ni-CeO₂/SBA-15-V. All samples showed type IV isotherm with a hysteresis loop in the relative pressure 0.65–0.80, confirming the uniform mesoporous structure in these samples. As shown in Figure 1b, the pore size of all samples was 6.5 nm, indicating the addition of Ni and CeO₂ to SBA-15 without changing the mesoporous structures. As listed in Table 1, the Brunauer-Emmett-Teller (BET) surface area and pore volume decreased after the addition of NiO and CeO₂ into SBA-15.

![Figure 1](image_url)

**Figure 1.** (a) N₂ adsorption-desorption isotherms; (b) pore size distributions of SBA-15, Ni-CeO₂/SBA-15-air, and Ni-CeO₂/SBA-15-V.

| Samples                  | \( S_{\text{BET}} \) (m²/g) | Pore Volume (cm³/g) | Average Pore Diameter (nm) |
|--------------------------|------------------------------|---------------------|----------------------------|
| SBA-15                   | 690                          | 1.12                | 6.56                       |
| Ni-CeO₂/SBA-15-air       | 433                          | 0.83                | 6.54                       |
| Ni-CeO₂/SBA-15-V         | 492                          | 0.83                | 6.56                       |

Table 1. Textural properties of SBA-15, Ni-CeO₂/SBA-15-air, and Ni-CeO₂/SBA-15-V.

Figure 2 shows the XRD patterns of the catalysts. The reduced catalysts were reduced in a 20% H₂/Ar stream (50 mL/min) for 40 min at 450 °C. As shown in Figure 2a, all the small-angle XRD patterns showed three typical peaks of mesoporous SBA-15 [27]. The result demonstrates that the mesoporous structure of SBA-15 was maintained after loading NiO and CeO₂. Figure 2b shows the wide-angle XRD patterns of fresh and reduced catalysts. Compared with Ni-CeO₂/SBA-15-air, Ni-CeO₂/SBA-15-V exhibited obviously wider reflection peaks of NiO and CeO₂, indicating that vacuum treatment can improve the dispersion of NiO and CeO₂. After reduction, the reflection peaks
of Ni in Ni-CeO₂/SBA-15-V were also wider than those in Ni-CeO₂/SBA-15-air, confirming that the Ni particle was smaller in the Ni-CeO₂/SBA-15-V catalyst after reduction. As listed in Table 2, the crystal sizes of Ni and CeO₂ in reduced Ni-CeO₂/SBA-15-V were 8.5 and 4.2 nm, which were obviously smaller than those in reduced Ni-CeO₂/SBA-15-air. The XRD results indicate that vacuum thermal treatment can improve Ni and CeO₂ dispersion on SBA-15 support.

Figure 3 shows TEM images of the catalysts. For all catalysts, a well-ordered mesoporous structure could be observed. For fresh Ni-CeO₂/SBA-15-air catalyst, most particles were 20–100 nm in size, which were larger than the pore size of SBA-15 and dispersed on the outside surface of SBA-15 (Figure 3a), and NiO particles larger than 20 nm could be clearly observed in Figure 3b. On the contrary, for fresh Ni-CeO₂/SBA-15-V (Figure 3c), lots of particles were aligned within the mesoporous channels of SBA-15, indicating this part of particles located in the pore channels. For fresh Ni-CeO₂/SBA-15-V, small particles were dispersed on the outside surface or in the nanochannel of SBA-15. NiO particles smaller than 10 nm in contact with CeO₂ particles were observed (Figure 3f).

Figure 4 shows H₂-TPR profiles of the catalysts. For Ni-CeO₂/SBA-15-air, the sharp peak located at around 420 °C was attributed to the reduction of large bulk NiO species with no or very little interaction with SiO₂ support [19]. For Ni-CeO₂/SBA-15-V, a reduction peak between 300 °C and 600 °C was observed. A reduction peak beyond 500 °C corresponds to the reduction of small NiO particles strongly interacting with SiO₂ support [28–33]. The result indicates that there is a strong interaction between Ni and SiO₂ in Ni-CeO₂/SBA-15-V catalyst.

| Catalysts                     | CeO₂ Crystal Size (nm) | NiO Crystal Size (nm) | Ni Crystal Size (nm) | Ni Dispersion (%) |
|-------------------------------|------------------------|-----------------------|----------------------|------------------|
| Ni-CeO₂/SBA-15-air (fresh)    | 7.3                    | 42.1                  | -                    | -                |
| Ni-CeO₂/SBA-15-air (reduced)  | 7.1                    | -                     | 26.1                 | 3.7              |
| Ni-CeO₂/SBA-15-V (fresh)      | 4.6                    | 10.8                  | -                    | -                |
| Ni-CeO₂/SBA-15-V (reduced)    | 4.2                    | -                     | 8.5                  | 11.4             |

* Determined by Scherrer’s equation from the (111) plane of CeO₂ in X-ray diffraction (XRD) patterns; b determined by Scherrer’s equation from the (200) plane of NiO in XRD patterns; c determined by Scherrer’s equation from the (111) plane of Ni in XRD patterns; d calculated as (97.1 nm)/(particle size of Ni(nm)).

Figure 2. (a) Small-angle and (b) wide-angle X-ray diffraction (XRD) patterns of SBA-15, Ni-CeO₂/SBA-15-air, and Ni-CeO₂/SBA-15-V catalysts.

Table 2. Physical properties of catalysts.
3.2. Catalytic Performance

The CO$_2$ conversion and CH$_4$ selectivity of the catalysts for CO$_2$ methanation reaction are shown in Figure 5. Compared with the Ni-CeO$_2$/SBA-15-air catalyst, the Ni-CeO$_2$/SBA-15-V catalyst exhibited obviously higher CO$_2$ conversion and CH$_4$ selectivity. For the Ni-CeO$_2$/SBA-15-V catalyst,
CO\textsubscript{2} conversion and CH\textsubscript{4} selectivity at 400 °C were 68.8% and 99.0%, respectively. CH\textsubscript{4} selectivity of the Ni-CeO\textsubscript{2}/SBA-15-V catalyst decreased to 97.1% at 450 °C. This is because reverse water gas shift reaction (CO\textsubscript{2} + H\textsubscript{2} \rightarrow CO + H\textsubscript{2}O) is an endothermic reaction, and high temperatures are more conducive to producing CO and decreasing CH\textsubscript{4} selectivity. The result shows that the Ni-CeO\textsubscript{2}/SBA-15-V catalyst was active and selective for the CO\textsubscript{2} methanation reaction. It should be noted that the Ni-CeO\textsubscript{2}/SBA-15-air catalyst produced a large amount of CO even at low temperature. This may be due to that the large Ni particle in the Ni-CeO\textsubscript{2}/SBA-15-air catalyst was active for both CO\textsubscript{2} methanation and reverse water-gas shift reaction, while small Ni particle in contact with CeO\textsubscript{2} in Ni-CeO\textsubscript{2}/SBA-15-V catalyst was more selective for CO\textsubscript{2} methanation.

![Figure 5. (a) CO\textsubscript{2} conversion and (b) CH\textsubscript{4} selectivity of Ni-CeO\textsubscript{2}/SBA-15-air and Ni-CeO\textsubscript{2}/SBA-15-V catalysts in methanation reaction.](image1)

Figure 5 shows the stability test of Ni-CeO\textsubscript{2}/SBA-15-V catalyst at 350 °C and 400 °C. The equilibrium conversions are also shown in Figure 6. The CO\textsubscript{2} conversions of Ni-CeO\textsubscript{2}/SBA-15-V catalyst had a little decrease in initial 10 h on stream, then the catalyst exhibited stable catalytic performance for 50 h at 350 °C and 400 °C. After reaction for 60 h, the CO\textsubscript{2} conversion at 350 °C and 400 °C stayed at around 49.5% and 62.5%, respectively. It is worth noticing that, during the stability test at 400 °C, CH\textsubscript{4} selectivity achieved over the Ni-CeO\textsubscript{2}/SBA-15-air catalyst remained almost constant after 10 h on stream approaching values higher than 97%.

![Figure 6. Stability of the Ni-CeO\textsubscript{2}/SBA-15-V catalyst in methanation reaction at (a) 350 °C and (b) 400 °C.](image2)

Among Ni-based methanation catalysts, Ni/Al\textsubscript{2}O\textsubscript{3} catalyst is the most widely investigated and used due to its low costs and availability. Table 3 shows the CO\textsubscript{2} conversion and selectivity of reported Ni/Al\textsubscript{2}O\textsubscript{3} catalyst at 250 and 300 °C. Compared with the Ni/Al\textsubscript{2}O\textsubscript{3} catalyst, the CO\textsubscript{2} conversion of Ni-CeO\textsubscript{2}/SBA-15-V was obviously higher.
was characterized by TG, XRD, and TEM. TG curve was shown in Figure 8a. There was a weight decrease below 100 °C and a weight increase in the temperature region of 180–400 °C. The weight decrease should be due to the desorption of adsorbed water and the weight increase due to the oxidation of Ni to NiO. There was no obvious weight loss above 100 °C, indicating no carbon deposition was formed on the spent Ni-CeO2/SBA-15-V catalyst. XRD patterns of reduced and spent Ni-CeO2/SBA-15-V catalysts were shown in Figure 8b. Determined by Scherrer’s equation from the (111) plane of Ni in XRD patterns, Ni crystal sizes were 8.5 and 9.1 nm for reduced and spent Ni-CeO2/SBA-15-V catalysts, respectively. XRD results indicate that the Ni particle size was slightly increased during the stability test. As shown in Figure 8c, TEM image also showed that no obviously sintering of Ni particle occurred. The TG, XRD, and TEM results indicate the slight decrease of activity and selectivity during stability test was due to the slight increase of Ni particle size.

The present work clearly demonstrates that the Ni-CeO2/SBA-15-V catalyst possessed superior activity and selectivity and exhibited excellent stability during the CO2 methanation reaction. The XRD and TEM results prove that the vacuum-thermal treated Ni-CeO2/SBA-15-V catalyst had higher Ni dispersion and smaller Ni particle size than the Ni-CeO2/SBA-15-air catalyst thermal treated in air. Based on the characterization and catalyst performance results, it can be concluded that good activity and selectivity of Ni-CeO2/SBA-15-V catalyst can be attributed to the small Ni particle size in contact with CeO2.

For comparison with Ni-CeO2/SBA-15-V catalyst, the stability of Ni-CeO2/Al2O3 catalyst prepared with a conventional impregnation method was tested at 350 °C. As shown in Figure 7, the CO2 conversion and CH4 selectivity of Ni-CeO2/Al2O3 catalyst were obviously lower than that of Ni-CeO2/SBA-15-V catalyst.

Carbon deposition and sintering of metal particle are the main reasons for the catalyst deactivation during CO2 methanation [10,35,36]. The spent Ni-CeO2/SBA-15-V catalyst after stability test at 400 °C was characterized by TG, XRD, and TEM. TG curve was shown in Figure 8a. There was a weight decrease below 100 °C and a weight increase in the temperature region of 180–400 °C. The weight decrease should be due to the desorption of adsorbed water and the weight increase due to the oxidation metal Ni to NiO. There was no obvious weight loss above 100 °C, indicating no obviously carbon deposition was formed on the spent Ni-CeO2/SBA-15-V catalyst. XRD patterns of reduced and spent Ni-CeO2/SBA-15-V catalysts were shown in Figure 8b. Determined by Scherrer’s equation from the (111) plane of Ni in XRD patterns, Ni crystal sizes were 8.5 and 9.1 nm for reduced and spent Ni-CeO2/SBA-15-V catalysts, respectively. XRD results indicate that the Ni particle size was slightly increased during the stability test. As shown in Figure 8c, TEM image also showed that no obviously sintering of Ni particle occurred. The TG, XRD, and TEM results indicate the slight decrease of activity and selectivity during stability test was due to the slight increase of Ni particle size.

| Catalysts                  | Ni Content (wt %) | T (°C) | CO2 Conversion (%) | CH4 Selectivity (%) | Ref.          |
|---------------------------|-------------------|--------|-------------------|---------------------|---------------|
| Ni-CeO2/SBA-15-V          | 10%               | 250    | 10.2              | 100                 | This work a   |
|                           |                   | 300    | 35.4              | 99.3                | This work a   |
| Ni/Al2O3                  | 14%               | 250    | 1                 | 100                 | [34] b        |
|                           |                   | 300    | 6                 | 100                 | [34] b        |

\[ a \] Weight hourly space velocity (WHSV) = 60,000 mL g\(^{-1}\) h\(^{-1}\), CO\(_2\)/H\(_2\)/Ar = 1/4/5; \[ b \] Gas hourly space velocity (GHSV) = 52,300 h\(^{-1}\), CO\(_2\)/H\(_2\) = 1/5.

Figure 7. Stability of Ni-CeO2/Al2O3 catalysts in methanation reaction at 350 °C.
was mainly attributed to higher Ni and CeO$_2$ dispersion. Vacuum thermal treatment exhibited better activity and selectivity than the Ni-CeO$_2$/SBA-15-air catalyst thermal treated in air. The excellent catalytic performance of the Ni-CeO$_2$/SBA-15-V catalyst after stability test at 400 °C proves that the vacuum-thermal treated Ni-CeO$_2$/SBA-15-V catalyst had higher Ni dispersion and smaller Ni particle size than the Ni-CeO$_2$/SBA-15-air catalyst thermal treated in air. Based on the characterization and catalyst performance results, it can be concluded that good activity and stability of Co catalysts for CO$_2$ methanation. Appl. Catal. B 2018, 220, 397–408. [CrossRef]

4. Conclusions

The effects of vacuum thermal treatment on Ni-CeO$_2$/SBA-15 catalysts were investigated in this paper. Compared with thermal treatment in air, vacuum thermal treatment can improve the dispersion of Ni and CeO$_2$ in the Ni-CeO$_2$/SBA-15 catalyst. The Ni-CeO$_2$/SBA-15-V catalyst with vacuum thermal treatment exhibited better activity and selectivity than the Ni-CeO$_2$/SBA-15-air catalyst thermal treated in air. The excellent catalytic performance of the Ni-CeO$_2$/SBA-15-V catalyst was mainly attributed to higher Ni and CeO$_2$ dispersion.

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