Preparation of Mesoporous Silica by Nonionic Surfactant Micelle–Templated Gelation of Na$_2$SiO$_3$ and H$_2$SiF$_6$ and Application as a Catalyst Carrier for the Partial Oxidation of CH$_4$

Kyeong-Won Park$^1$, Jin-Young Kim$^2$, Ho-Joon Seo$^2$ & Oh-Yun Kwon$^2$

Mesoporous silica (MSPN12) was prepared by nonionic surfactant micelle–templated gelation of sodium silicate (Na$_2$SiO$_3$) and fluorosilicic acid (H$_2$SiF$_6$) in aqueous solution, characterized by a range of instrumental techniques, and tested as a support for Ni and Rh catalysts in the partial oxidation of methane (POM). Calcined and sintered MSPN12 exhibited well-defined $d_{001}$-spacings (3.5–4.39 nm), narrow pore distributions (2.4–3.1 nm), and large specific surface areas (552–1,246 m$^2$ g$^{-1}$), and was found to be highly thermally stable. Microscopic imaging revealed that MSPN12 comprised spherical particles with a uniform diameter of ~0.7 µm, with each particle featuring firm and regular honeycomb-type pores. MSPN12-loaded Ni and Rh maintained stable POM activity at 700 °C during almost 100 h on stream, which were comparable to those for the commercial Rh(5)/Al$_2$O$_3$ catalyst in terms of methane conversion and H$_2$ formation selectivity. Thus, the combination of structural stability and favorable physicochemical properties resulted in good POM performance.

Micro- and mesoporous materials are often used as catalyst supports, adsorbents, and ion exchangers. For example, zeolites are widely applied as adsorbents and catalyst supports because of their acidic properties and ion-exchange ability. However, the use of zeolites as catalysts for heavy oil cracking is hindered by their small pore size (< 1 nm). To overcome this problem, researchers at Mobil Oil Corporation developed MCM-41 and MCM-48. Since then, numerous MCM-41–related materials have been synthesized and applied in the field of nanochemistry as catalysts, adsorbents, and catalyst carriers. Typically, these materials are prepared using surfactant micelle–based templating, and their pore properties can therefore be altered by changing micelle structure and morphology, e.g., via variation of surfactant type, molecular structure, concentration, and additives.

MCM-41–type materials synthesized by cationic surfactant micelle–based templating exhibit a very regular and uniform pore structure and are therefore expected to be superior catalyst carriers. However, low pore wall thickness results in the easy collapse of these pore structures under hydrothermal conditions or at temperatures above 700 °C, which can be overcome through the use of nonionic surfactant micelle–based templates. Nonionic surfactants commonly contain ethylene oxide chains that can participate in dipole interactions with metal hydroxide ions. When these ions are stabilized by dipole interactions and solidified via condensation, the pore walls of the resulting mesoporous materials are expected to be thicker than the ethylene oxide chain length. When such micelle templates are calcined, mesopores form in the alkyl chain–containing core portion, while micropores are formed in ethylene oxide chain–containing pore walls. In this case, microporous pore walls are expected to be very thick, and the corresponding supported catalysts are expected to have large specific surface areas.

---

$^1$Department of Chemistry and Faculty of General Education, Gyeongsang National University, Jinju, 52828, Republic of Korea. $^2$Department of Chemical and Biomolecular Engineering, Chonnam National University, Yosu, Chonnam, 59626, Republic of Korea. Correspondence and requests for materials should be addressed to K.-W.P. (email: nano2k@nate.com) or O.-Y.K. (email: oykwon@jnu.ac.kr)
areas and high thermal stabilities. However, despite these advantages, the synthesis of catalysts via nonionic surfactant–based templating and their applications have been underexplored.[9-13].

Generally, high-purity mesoporous silica (MS) is prepared from silica gel or tetraethylorthosilicate. However, recent studies utilized sodium silicate (Na₂SiO₃) or fluorosilicic acid (H₂SiF₆) as precursors to minimize problems such as waste solution treatment, which is time-consuming, expensive, and requires the use of non-aqueous solutions.

Herein, MS synthesized by simultaneous gelation of Na₂SiO₃ and H₂SiF₆ was tested as a catalyst support for the partial oxidation of methane (POM). In this synthesis, H₂SiF₆ acted as an acid and silica source, additionally supplying fluoride anions to promote mineralization. As H₂SiF₆ can easily dissolve metal oxides and organo-metallic compounds, it allows one to introduce various metals in the pore walls of silica skeletons. Additionally, H₂SiF₆ is a very cheap raw material, since it is a by-product of phosphate fertilizer production. The reaction between H₂SiF₆ and Na₂SiO₃ was completed within several seconds, which allowed the synthesis of MS to be carried out as a continuous process. MS synthesized in this manner featured large surface area and high thermal stability, and was concluded to be a superior catalyst support for high-temperature reactions.

In recent studies, hydrogen and CO₂ are highlighted from economic and environmental aspects. Fuel cells are attractive power supplies for versatile applications of hydrogen. In general, hydrogen is obtained by the steam reforming of methane[14]; recently, renewable oxygenated hydrocarbons such as methanol, ethanol, dimethyl ether (DME), and glycerol have also been used[15,16]. CO₂ methanation[17,18] and oxidative dehydrogenation of ethane using CO₂[19] are promising routes for converting CO₂ into fuels and chemicals toward CO₂ emission control.

In this study, we tested MS as the catalyst carrier for the partial oxidation of methane (POM).

Materials and Methods
Materials. Na₂SiO₃ (35–38 wt% SiO₂, Kanto Chemical Co., Japan), H₂SiF₆ (35 wt%, Alfa Aesar, England), and polyoxyethylene(12) nonylphenol ether (PN12; Aldrich, USA) were used for the preparation of mesoporous silica. Ni(NO₃)₂·6H₂O (Aldrich, USA), RhCl₃·xH₂O (Aldrich, USA), commercial Rh(5)/Al₂O₃ (Aldrich, metal loading = 5 wt%), and ethanol (Aldrich, USA) were used for catalyst preparation.

Preparation of mesoporous silica (MSPN12). Aqueous solutions of Na₂SiO₃ (10 wt%), PN12 (3.0 wt%), and H₂SiF₆ (6.0 wt%) were mixed to achieve a Na₂SiO₃·H₂SiF₆·H₂O ratio (w/w) of 1:20:3.75:459. First, the surfactant solution was added to the Na₂SiO₃ solution, and the resulting dispersion was homogeneously mixed and stirred at 400 rpm for 0.5 h in a constant-temperature reactor at 50 °C. At this point, pH was measured as 11. Subsequently, the H₂SiF₆ solution was added in one portion to the mixed solution maintained at 50 °C, which resulted in the formation of a white precipitate within 5 s and a decrease of pH to 5.0. The obtained precipitate was filtered, dried at 60 °C, and calcined in air at 500 °C for 3 h to remove the surfactant template and afford the MSPN12 molecular sieve with a constant pore size. The thermal stability of MSPN12 was tested by additional 3-h sintering at 700, 800, and 900 °C.

Preparation of catalysts. Ni(5)/MSPN12 and Rh(5)/MSPN12 catalysts (metal loading = 5 wt% each) were prepared as follows. MSPN12 (1.0 g) was dispersed in 5 mL of ethanolic nickel nitrate (containing 0.06 g of Ni(NO₃)₂·6H₂O) or rhodium nitrate (containing 0.01 g of RhCl₃·xH₂O) solutions upon 5-h stirring, and the resulting dispersions were evaporated, dried for 24 h at 100 °C, and annealed for 5 h at 500 °C in an electric furnace (Eyela TMF-1000) in air. Ethanol was found to be superior to water, allowing one to realize a more uniform dispersion of metals at the gallery surface. Commercial Rh(5)/Al₂O₃ was tested together with the above catalysts for comparison.

Catalytic reaction. The catalytic reaction was carried out under atmospheric pressure in a fixed-bed flow reactor comprising a quartz tube (10 mm inner diameter) and catalyst powder (0.05 g) held on quartz wool. The CH₄/O₂ molar ratio, temperature, pressure, and the gas hourly space velocity (GHSV) of the reactant gas equaled 2, 700 °C, 1 atm, and 7.32 × 10⁴ mL g⁻¹ cat⁻¹ h⁻¹, respectively. The reactor was maintained at the desired temperature with an accuracy of ±1 °C using a proportional-integral-derivative controller and a K-type thermocouple located on the catalyst. The reactants were purged using a pressure regulator attached to the corresponding gas cylinders, and the composition of the reactant mixture was controlled using mass flow meters. The effluent was analyzed using an online gas chromatograph (Shimadzu Co., Model 14B, Japan) equipped with a thermal conductivity detector and Porapak Q and 5 Å molecular sieve columns arranged in parallel. The fresh catalyst was reduced in a flow of hydrogen (20 mL min⁻¹) at 500 °C for 5 h, and the temperature was subsequently increased to 700 °C at a rate of 10 °C min⁻¹ before exposure to reactant gases.

Catalyst characterization. X-ray diffraction (XRD) patterns were recorded on a Bruker D8 Advance diffractometer (Cu Kα radiation, λ = 1.5406 Å; 40 kV, 40 mA). Scanning electron microscopy (SEM) imaging was carried out using a JEOL JSM-840A microscope. Samples intended for SEM imaging were stuck onto adhesive tape, sputter-coated with gold, and imaged. The transmission electron micrographs were obtained with a JEOL JEM-200 CX microscope operated at 200 kV, using the thin-section technique. The powder samples were embedded in epoxy resin and sectioned with an ultra-microtome. N₂ adsorption/desorption isotherms were recorded by a Micromeritics ASAP 2020 instrument at −196 °C for samples outgassed in vacuum for 4 h at 300 °C. Specific surface areas were determined using the BET method[20], and pore size distributions were determined using N₂ adsorption/desorption data and the BJH method[21].
Results and Discussion

The gelation of Na<sub>2</sub>SiO<sub>3</sub> with H<sub>2</sub>SiF<sub>6</sub> in the presence of a surfactant micelle template was completed within several seconds, affording uniform spherical particles. Figure 1 shows the XRD patterns of MSPN12 sintered for 3 h at 700, 800, and 900 °C after 3-h calcination at 500 °C. MSPN12 calcined at 500 °C showed a well-developed peak corresponding to a d-spacing of 4.39 nm, which indicated that surfactant micelles served as templates to form homogeneous pore. The intensity and half-height width of this peak decreased with increasing sintering temperature, reflecting the unclear distinction between pores and pore walls due to the concomitant partial collapse of the pore structure. However, the peak was maintained at 900 °C, which indicated the high thermal stability of MSPN12. The employed nonionic surfactant, PN12, comprised a hydrophilic polyoxyethylene chain [(CH<sub>2</sub>CH<sub>2</sub>O)<sub>12</sub>], which was stabilized by dipole interactions with monomers such as Si(OH)<sub>4</sub>, SiO(OH)<sub>3</sub><sup>−</sup>, and SiO<sub>2</sub>(OH)<sub>2</sub><sup>2−</sup> and solidified via the polycondensation reaction. Consequently, solidification of the monomers between the polyoxyethylene chains results in thicker pore walls than the polyoxyethylene chain length, thus ensuring high thermal stability.

In Na<sub>2</sub>SiO<sub>3</sub> solutions, Si(OH)<sub>4</sub> is the predominant species at pH ≤ 7, while in solutions with pH > 9, anionic species such as SiO(OH)<sub>3</sub><sup>−</sup> and SiO<sub>2</sub>(OH)<sub>2</sub><sup>2−</sup> are converted into grains through hydrolytic condensation reactions. However, in aqueous nonionic surfactant solutions of 3 ≤ pH ≤ 9, Si(OH)<sub>4</sub> precipitates into SiO<sub>2</sub> within several seconds. Here, Si(OH)<sub>4</sub> groups form hydrogen bonds with the oxygens of the polyoxyethylene chains.
which promotes the formation of SiO(OH)$_3$ anions to induce SiO$_2$ precipitation. SiF$_6^{2-}$ ions in aqueous cationic or nonionic surfactant solutions also precipitate into SiO$_2$, within several seconds (even in acidic solutions with pH 3–4)\textsuperscript{15,17,27} which is ascribed to the mineralization-promoting action of F$^-$ ions, similar to that of OH$^-$ ions. Therefore, the gelation of Na$_2$SiO$_3$ with H$_2$SiF$_6$, in the aqueous surfactant solution was promoted by the mineralizing effect of the F$^-$ ions and the electrostatic or dipolar interactions between the surfactant micelles and silica.

| Temperature (°C) | $d_{100}$ (nm) | BET surface area (m$^2$ g$^{-1}$) | BJH pore diameter (nm) adsorption |
|------------------|----------------|----------------------------------|---------------------------------|
| 500              | 4.39           | 1.246                            | 3.1                             |
| 700              | 4.05           | 1.088                            | 2.7                             |
| 800              | 3.82           | 882                              | 2.5                             |
| 900              | 3.50           | 552                              | 2.4                             |

Table 1. Physical properties of MSPN12 calcined and sintered at various temperatures.
monomers (Si(OH)₄, SiO(OH)₂⁻, SiO₂(OH)₃⁻, etc.). Silva and Pastore reported that a more uniform pore structure of mesoporous silica is formed in the presence of fluoride ions and ascribed this behavior to the promotional effect of these ions on the solidification of MSPN12 pore walls, additionally demonstrating that this effect results in enhanced thermal stability.

Figure 2 shows typical SEM and TEM images of MSPN12, revealing that this material comprised well-separated spherical particles with a uniform diameter of ~0.7 μm, which was ascribed to the fast rate of their formation. Generally, large particles grow around seeds when the rate of crystal seed formation is slow, whereas at higher seed formation rates, the number of these seeds increases so much that the growth of particles stops, which affords smaller particles with a uniform size. In contrast to mesoporous silica obtained using nonionic surfactants, which was reported to exhibit irregular wormhole-type pores opening in all directions, MSPN12 featured pores with a firm and regular honeycomb-like structure.

Figure 3 shows the N₂ adsorption isotherms of MSPN12 sintered at various temperatures, demonstrating a typical steep increase with mesopore filling at a relative pressure of 0.3–0.4 for all samples.

Table 1 lists the specific surface areas obtained using the BET equation, showing that the largest area of 1,246 m² g⁻¹ was observed for MSPN12 calcined at 500 °C. This finding was attributed to an increase of micropore surface area due to the effect of polyethylene oxide chains. Upon calcination of micelle templates, mesopores formed in the alkyl chain–containing core portion, while micropores formed in polyethylene oxide chain–containing pore walls, and the resulting mesoporous materials consequently exhibited a considerably large specific surface area. The increase of sintering temperature from 500 to 900 °C brought about a large decrease of surface area.
area from 1,246 to 552 m² g⁻¹ due to the thermally induced collapse of micropores. This trend was well reflected in the pore size distributions of MSPN12 (Fig. 4).

Figure 5 schematically illustrates the synthesis of MS by nonionic surfactant micelle–based templating, demonstrating that the thick and solid nature of pore walls was caused by the presence of polyethylene oxide chains. Monomers produced in the reaction of Na₂SiO₃ and H₂SiF₆ (e.g., Si(OH)₄, SiO₂(OH)₃⁻, and SiO(OH)₂⁻) were stabilized by dipolar interactions between polyethylene oxide chains at the micelle surface. During the subsequent condensation reaction, this stabilization led to the formation of a thick SiO₂ skeleton in large spaces between micelles, which endowed MSPN12 with a large surface area and high thermal stability.

Finally, MSPN12 was examined as a catalyst carrier for POM, with CH₄ conversions achieved for Ni(5)/MSPN12, Rh(5)/MSPN12, and Rh(5)/Al₂O₃ shown in Fig. 6.

Ni(5)/MSPN12 and Rh(5)/MSPN12 remained stable for almost 100 h on stream, and CH₄ conversion (>90%) over these catalysts exceeded that (>80%) over the commercial catalyst. The trends of H₂ yield, shown in Fig. 7, were similar to those observed for methane conversion, and the high activity and stability of Ni(5)/MSPN12 were consequently concluded to hold great promise for practical applications.
H₂ yields of nearly 90% were observed for Ni(5)/MSPN12 and Rh(5)/MSPN12, whereas the corresponding yields of (CH₃)₂ were low, reflecting minimal carbon deposition. Generally, carbon deposition on the catalyst surface results in a decrease of catalytic activity. However, carbon deposition did not disturb the stable performance of MSPN12 catalysts.

In particular, metal particles (Fig. 8) were uniformly dispersed within the mesopores owing to the confinement effect. This uniform dispersion of the metal in the mesopores also contributes to the prolonged maintenance of their catalytic activity. Thus, the obtained results indicated that the large surface area, firm and ordered pore structure, and high thermal stability of MSPN12 contributed to the good POM performance of MSPN12-supported catalysts.

Conclusion
Herein, mesoporous silica was synthesized through the gelation of Na₂SiO₃ and H₂SiF₆ in water templated by nonionic surfactant micelles. The above reaction was completed within 5s and afforded a precipitate comprising well-separated particles with a uniform diameter of ~0.7 μm and a regular honeycomb-like pore structure. Structural analysis of this precipitate allowed it to be classified as a mesoporous molecular sieve. The corresponding specific surface areas were in the range of 552–1,246 m² g⁻¹ and depended on sintering temperature, while pore size distributions featured maxima at 2.4–3.1 nm. The synthesized mesoporous silica was shown to be thermally stable up to 900°C.

Ni- and Rh-impregnated mesoporous silica catalysts remained stable for almost 100 h on stream under the conditions of partial methane oxidation and achieved methane conversions (>90%) exceeding those (>80%) obtained over a commercial Rh(5)/Al₂O₃ catalyst, with a similar trend observed for hydrogen yield. The unique properties of mesoporous silica such as its large surface area, ordered pore structure, and high thermal stability were concluded to contribute to the good performance of mesoporous silica–supported catalysts for partial methane oxidation and hold great promise for practical applications.

References
1. Ang, Y. et al. Uniform nanosize mesopores with large secondary pore architecture. Chem. Mater. 18, 1861–1866 (2006).
2. Krege, C. T., Leonowicz, M. E., Roth, W. J., Vartuli, J. C. & Beck, J. S. Ordered mesoporous molecular sieves synthesized by a liquid-crystal template mechanism. Nature 359, 710–712 (1992).
3. Beck, J. S. et al. A new family of mesoporous molecular sieves prepared with liquid crystal templates. J. Am. Chem. Soc. 114, 10834–10843 (1992).
4. Beck, J. S. et al. Molecular or supramolecular templating: defining the role of surfactant chemistry in the formation of microporous and mesoporous molecular sieves. Chem. Mater. 6, 1816–1821 (1994).
5. Hsu, Q., Leon, R., Petroff, P. M. & Stucky, G. D. Mesostructure design with gemini surfactants: supercage formation in a three-dimensional hexagonal array. Science 268, 1324–1327 (1995).
6. Zhao, D. et al. Triblock copolymer syntheses of mesoporous silica with periodic 50 to 300 angstrom pores. Science 279, 548–552 (1998).
7. Tanev, P. T. & Pinnavaia, T. J. A neutral templating route to mesoporous molecular sieves. Science 267, 865–867 (1995).
8. Ryoo, R., Kim, J. M., Ko, C. H. & Shin, C. H. Disordered molecular sieve with branched mesoporous channel network. J. Phys. Chem. 100, 17718–17721 (1996).
9. Inagaki, S., Ogata, S., Goto, Y. & Fukushima, Y. Mesoporous materials derived from layered silicates and the adsorption properties. Stud. Surf. Sci. Catal. 117, 65–76 (1998).
10. Joo, S. H. et al. Ordered nanoporous arrays of carbon supporting high dispersions of platinum nanoparticles. Nature 412, 169–172 (2001).
11. Lee, J., Sohn, K. & Hyeon, T. Fabrication of novel mesocellular carbon foams with uniform ultralarge mesopores. J. Am. Chem. Soc. 123, 5146–5147 (2001).
12. Bagshaw, S. A., Prouzet, E. & Pinnavaia, T. J. Templating of mesoporous molecular sieves by nonionic polyethylene oxide surfactants. Science 269, 1242–1244 (1995).
13. Sierra, L., Lopez, R., Gil, H. & Guth, J. L. Synthesis of mesoporous silica from sodium silica solutions and a poly(ethylene oxide)-based surfactant. Adv. Mater. 11, 307–311 (1999).
14. Venugopal, E., Aswal, V. K. & Kumaraswamy, G. Nanoparticle size controls aggregation in lamellar nonionic surfactant mesophase. Langmuir 29, 9643–9650 (2013).
15. Kwon, O. Y., Kim, S. W. & Choi, S. W. Synthesis of mesoporous molecular sieves: hydrolysis of H₂SiF₆ by a non-ionic polyethyleneoxide surfactant template. Microporous Mesoporous Mater. 27, 255–259 (1999).
16. Luan, Z., He, H., Zhou, W. & Klinowski, J. Transformation of lamellar silicate into the mesoporous molecular sieve MCM-41. J. Chem. Soc., Faraday Trans. 94, 979–983 (1998).
17. Jeong, S. Y., Suh, J. K., Lee, J. M. & Kwon, O. Y. Preparation of silica-based mesoporous materials from fluorosilicon compounds: gelation of H₂SiF₆ in ammonia surfactant solution. J. Colloid Interface Sci. 192, 156–161 (1997).
18. Engler, B. C., Ludeng, R. & Holmsen, A. Appl. Catal. A: General 346, 1–27 (2008).
19. Ma, K. et al. ChemCatChem 10, 4010–4017 (2018).
20. Ma, K. et al. Chem. Sci. 10, 2578–2584 (2019).
21. Ma, Y. et al. Chem. Eng. Sci. 194, 10–21 (2019).
22. Lu, Z. et al. ChemCatChem 10, 720–724 (2018).
23. Zhang, R. et al. ACS Catal. 8, 9280–9286 (2018).
24. Brunauer, S., Emmett, P. H. & Teller, E. Adsorption of gases in multimolecular layers. J. Am. Chem. Soc. 60, 309–319 (1938).
25. Barrett, E. P., Joyner, L. G. & Halenda, P. P. The determination of pore volume and area distributions in porous substances. I. Computations from nitrogen isotherms. J. Am. Chem. Soc. 73, 373–380 (1951).
26. Iller, R. K. The Chemistry of Silica: Solubility, Polymerization, Colloid and Surface Properties, and Biochemistry (Wiley-Interscience, New York, USA 1979).
27. Kwon, O. Y., Jeong, S. Y., Suh, J. K., Choi, S. W. & Lee, J. M. Synthesis of mesoporous molecular sieves from fluorosilicon compound (H₂SiF₆). Bull. Korean Chem. Soc. 18, 663–665 (1997).
28. Silva, F. H. P. & Pastore, H. O. The syntheses of mesoporous molecular sieves in fluoride medium. Chem. Commun. 0, 833–834 (1996).
29. Zhang, S., Muratsugu, S., Ishiguro, N. & Tada, M. Ceria-doped Ni/SBA-16 catalysts for dry reforming of methane. ACS Catal. 3, 1855–1864 (2013).
Author Contributions
K.W.P. and J.Y.K. performed the experiments. SEM/TEM/XRD/BET and all figure drawing were by K.W.P. and H.J.S. carried out catalytic testing. K.W.P. and O.Y.K. wrote the manuscript writing.

Additional Information
Competing Interests: The authors declare no competing interests.

Publisher's note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit http://creativecommons.org/licenses/by/4.0/.

© The Author(s) 2019