Correlating the effect of preparation methods on the structural and magnetic properties, and reducibility of CuFe$_2$O$_4$ catalysts

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CuFe$_2$O$_4$ spinel oxide has attracted research interest because of its versatile practical applications, especially for catalysis. In this study, nanometre-sized CuFe$_2$O$_4$ particles were prepared by three different methods, including nanospace confinement in SBA-15, hard template removal, and sol–gel combustion. The relationship between structure, size, magnetic behaviour, and reducibility of the catalysts was further investigated by various advanced techniques. Samples prepared by impregnation and hard template removal show high surface area and small crystallite size with superparamagnetic behaviour. In contrast, the sol–gel sample exhibits ferromagnetic properties with a large crystallite size and low surface area. Although all samples present a tetragonal crystal structure, the distributions of Fe and Cu cations in tetrahedral and octahedral sites in the spinel structure are different. The reducibility results demonstrate that the supported CuFe$_2$O$_4$/SBA-15 shows the lowest reduction profile. These results could suggest that the synthesis method strongly affects the crystal properties and cation distribution in the spinel structure, microstructure, surface area and reducibility, which are among the most relevant physicochemical properties for the catalytic activity.

1. Introduction

Copper spinel oxides have long attracted attention of researchers because their structural property is suitable for use as precursors or model catalysts.1–4 The structure of copper spinel provided excellent performance in the degradation of oxy-hydrocarbons at relatively lower temperatures of 200–350 °C, compared to other types of Cu-based catalysts.5–8 A series of Cu-spinel species of CuMn$_2$O$_4$6, ZrO$_2$–CuZn(Al$_2$O$_3$)7, and CuFe$_2$O$_4$–SiO$_2$ catalysts6 have been investigated lately. Among them, CuFe$_2$O$_4$ is considered to be a promising catalyst for methanol steam reforming because of the abundance of Cu and Fe in nature and its active ability for hydrogen production. However, the CuFe$_2$O$_4$ spinel catalysts typically possessed low surface area and their copper particles became sintered after high temperature calcination, resulting in insufficient active sites. It is well known that the activity of CuFe$_2$O$_4$ spinel is greatly dependent on Cu particle size and its dispersion. Well dispersed Cu particles with a slow rate of sintering can promote high hydrogen production yield.

We recently examined CuFe$_2$O$_4$ catalysts prepared by a urea–nitrate combustion method for dimethyl ether steam reforming and found that these catalysts were highly active in spite of their low surface area.59 Nevertheless, the CuFe$_2$O$_4$ spinel in the form of nanopores and nanoparticles should be developed to provide a new structure with higher surface area, leading to better catalytic performance. A common method to meet this requirement is impregnating mesoporous silica powders with the desired salt precursors. A hard template for crystal growth of porous metal spinel oxide is also one of many interesting techniques for preparing high surface area materials, especially mesoporous metal oxides.60 Both methods have been used as a powerful strategy for producing crystalline nanoporous materials having a vast potential for applications in the
catalysis. Some advanced techniques as a block copolymer templates have been demonstrated for controlling the size, morphology, and shape of nanoparticles. To the best of our knowledge, however, there are a few studies reporting on the relationship between structure, magnetic behaviour, and reduction behaviour of CuFe$_2$O$_4$ spinel prepared with different methods. Thus, in this research, the relationships of those behaviours have been examined. We have expected that this investigation could provide an information for preparing a suitable catalyst, regarding to a significant role of preparation methods on structural behaviour and catalytic performance of CuFe$_2$O$_4$ catalysts.

2. Experimental

2.1 Catalyst preparation

2.1.1 Impregnation and template removal method. Mesoporous SBA-15 silica template was synthesized according to method described in ref. 17. Short descriptions of method were briefly discussed as follows; 4 g of pluronic P-123 (Sigma Aldrich) was added to 130 mL of H$_2$O and 20 mL of 1 M hydrochloric acid (HCl, 37%). Stirred until complete dissolution. Then, approximately 9 mL of tetraethylorthosilicate (TEOS, sigma Aldrich, 98%) was added to the solution and stirred for 7.5 h at 45 °C. After that, the sample was aged at 80 °C for 15.5 h without stirring, cooled down to room temperature, filtered, washed, and calcined at 550 °C for 6 h with a heating rate of 1 °C min$^{-1}$ in ambient atmosphere.

The CuFe$_2$O$_4$/SBA-15 was prepared by impregnation method using nitrate solutions of Cu and Fe with a 0.4 molar ratio. Firstly, the nitrate salts were dissolved in 0.8 mL of deionized (DI) water and placed in a vacuum oven at room temperature for 1 h. The powder of mesoporous SBA-15 was slowly added to the mixture to form a homogeneous solution (the total weight ratio of nitrate salts and mesoporous SBA-15 was 1:1). Later on, the mixture was placed on the hot plate at 70 °C for 1 h, ground and dried in vacuum oven overnight. After that, the red sample was calcined at 750 °C for 15 h with a heating rate of 1 °C min$^{-1}$ (the final metal loading was 27 wt. % and denoted as CuFe$_2$O$_4$/SBA-15).

2.1.2 Hard template removal method. The mesoporous SBA-15 template of CuFe$_2$O$_4$/SBA-15 was eliminated by washing in 50 mL of 2 M NaOH for three times. The residue metal oxides were further centrifuged and washed with 100 mL of 95% ethanol three times. The solid metal oxide was dried at 100 °C overnight. The red-brown product was obtained, denoted as CuFe$_2$O$_4$-TP.

2.1.3 Sol-gel combustion method. The sol-gel combustion with a citric acid complex method was employed in this work. Cu(NO$_3$)$_2$.2.5H$_2$O (Ajax, 98%) and Fe(NO$_3$)$_3$.9H$_2$O (Ajax, 98%) were dissolved in DI water and stirred continuously at 60 °C for 2 h. The mixture was added by excess amount of citric acid (Ajax, 98%) and kept stirring for 1 h. After that, the solution was heated up to 100 °C to evaporate water, then combusted at 200 °C until the reaction was completed. The combusted sample was calcined under air atmosphere at 750 °C for 15 h with a heating rate of 3 °C min$^{-1}$, denoted as CuFe$_2$O$_4$-SG.

2.2 Characterizations

The phase and crystallinity of samples were confirmed by powder X-ray diffraction (XRD; Bruker, D8 ADVANCE) with Cu K$_\alpha$ radiation at 40 kV and 40 mA, increment of 0.02° per second, step time of 0.5 s over the scan range of 10° < 2θ < 80°. The specific surface area and pore volume were determined using nitrogen adsorption–desorption isotherm technique (Bel, Belsorp-max) at 77 K. Each sample was outgassed overnight at 300 °C before measuring. The reduction profiles were carried out using temperature-programmed reduction (TPR; Quantachrome Instruments, CHEMIBET-Pulsar). About 25 mg of the calcined catalyst was heated under atmosphere of 5% H$_2$/Ar with 10 °C min$^{-1}$ heating rate and 30 mL min$^{-1}$ flow rate to final temperature of 800 °C. The morphologies of the catalysts were examined with transmission electron microscopy (TEM; JEOL, JEM-2100). The samples were dispersed in methanol media and then deposited on a Cu grid. Magnetic hysteresis loops were recorded at room temperature using a vibrating sample magnetometry (VSM; LAKE SHORE VSM 7404, Lake Shore Cryotronics, Inc., USA) with a maximum magnetic field of 10 kG. The geometry, local environment, and structural properties of CuFe$_2$O$_4$ were solved by X-ray absorption spectroscopy (XAS) technique. Transmission XAS spectra were collected at Fe and Cu K-edges at the SUT-NANOTEC-SLRI XAS beamline (BL5.2; electron energy of 1.2 GeV; bending magnet; beam current of 80–150 mA; 1.1 to 1.7 × 10$^{11}$ photon per s) of the Synchrotron Light Research Institute, Public Organization, Thailand. For the acquisition of all spectra, a Ge (220) double crystal monochromator with an energy resolution (ΔE/E) of 2 × 10$^{-4}$ was used. Energy calibrations were carried out using Fe foil (7112 eV) and Cu foil (8979 eV). In the X-ray absorption near edge structure (XANES) region, the step size was 0.02 eV with a second integration time. The extended X-ray absorption fine structure (EXAFS) region was set to scan to the maximum of 10k. The data reduction and EXAFS curve fitting were performed using the Athena and Artemis programs which are included in an IFEFFIT package.

2.3 Methanol steam reforming evaluation

The catalytic activity of the prepared catalysts was demonstrated for methanol steam reforming using a tubular reactor under atmospheric pressure. A 0.2 g of each catalyst was packed in a tubular quartz cell and sandwiched with quartz wool on both sides then placed inside the furnace. Prior to the experiments, the catalyst was reduced in 10% H$_2$ balanced with N$_2$ at 350 °C with a flow rate of 40 mL min$^{-1}$ for 3 h. After the reduction was completed, the temperature was maintained at 200 °C under N$_2$ flow. A mixture of methanol and steam (steam/carbon ratio = 2) was injected to a pre-heater at 150 °C to generate a gas feed stock using syringe pump with an injection rate of 100 mL min$^{-1}$. After that, the gas feed stock was introduced through the reactor by N$_2$ carrier gas with a flow rate of 100 mL min$^{-1}$. The reaction temperature was held in a range of 200–375 °C. The derived gaseous products in the supplied and reformed gases after experiment were collected at 3 °C using a condenser. The gas compositions were analyzed by online gas.
chromatography (GC) equipped with a thermal conductivity detector (VARIAN, CP-4900). A Poraplot U column was used to separate methanol, and CO₂. A capillary molecular sieve 5 Å column was applied for H₂, O₂, N₂, CH₄, and CO separations. The catalytic activity was evaluated by the H₂ production rate.

3. Results and discussion

3.1 Characterization of CuFe₂O₄ catalyst

3.1.1 Formation of CuFe₂O₄ spinel. The XRD patterns of all samples are displayed in Fig. 1. Fig. 1(A) shows low-angle XRD patterns of the SBA-15 and CuFe₂O₄/SBA-15 composite. It was found that the SBA-15 exhibited typical reflection of 2θ in the region between 0.8° and 2.5° corresponding to d-spacing of (100), (110), (200) and (210) ascribed to the d values (d defined as a distance between the empty pore space and the wall pore) of 10.2 nm, 6.0 nm, 5.3 nm, and 4.0 nm, respectively. These reflections possessed the 2D hexagonally structural pores belonging to P6/mmm space group of an ordered mesoporous silica. The ordered hexagonal tunnels remained after pores filling of metal precursors and further calcination under severe condition. It is noteworthy to mention that some spinel CuFe₂O₄ species possibly formed and grew inside the pores, leading to the reduction of SBA-15 diffraction peaks. The assumption was confirmed by the observed shrinkage of the spacing distances of (100), (110), and (200) which were equal to 9.8 nm, 5.9 nm, and 5.1 nm, respectively. Fig. 1(B) shows wide-angle XRD spectra of calcined samples with different preparation methods. Most of the diffraction peaks corresponded to the spinel phase of CuFe₂O₄, manifested by the sharp characteristic reflection planes. This lattice parameter for tetragonal spinel CuFe₂O₄ is in good agreement with that previously reported. In addition, XRD peaks of copper oxides and iron oxides were not detected in any samples. This would indicate that the synthesis methods were feasible to effectively prepare a pure CuFe₂O₄ phase. The catalyst prepared via sol–gel method showed sharp and intense reflection peaks. This was probably due to particle agglomeration during calcination. On the other hand, the XRD patterns of CuFe₂O₄/SBA-15 composite and CuFe₂O₄_TP show much broader peaks. Overlapping peaks of XRD profiles were evident and became dominant. This discernible feature indicated that small crystalline sizes without sintering of spinel oxides were formed, comparing to the sol–gel method. It should be noted that the SBA-15 has promoted and assisted the dispersion of CuFe₂O₄ on the surface. A closer look at these XRD peak profiles reveals a sharp 004 reflection which would suggest that the SBA-15 may favor the growth of CuFe₂O₄ crystal along one direction. The structure of spinel oxide (CuFe₂O₄_TP) was also maintained after leaching out of SBA-15. In addition, the SBA-15 did not present any diffraction patterns at high angles due to its amorphous nature of pore walls.

3.1.2 Morphology of CuFe₂O₄ spinel. To support our previous statements about pore filling, structure collapsing, and particle agglomeration, we have verified our inquisition by using TEM technique. Fig. 2 depicts the TEM images of SBA-15, CuFe₂O₄/SBA-15 and CuFe₂O₄ spinel synthesized with a sol–gel combustion and a template removal method. From the result, Fig. 2(A) displays a clear image of regular hexagonal pore array and empty mesopores SBA-15. The TEM image of CuFe₂O₄/SBA-15 (Fig. 2(B)) revealed the periodic nature of the pores, indicating that the structure of materials remained. Homogeneous dispersion of CuFe₂O₄ on SBA-15 was observed. The CuFe₂O₄ particles after leaching out the SBA-15 in NaOH solution exhibited small needle-like clusters with small particles dispersed on the surface (Fig. 2(C)). This one-dimensional growth of CuFe₂O₄ nanocrystals was in good agreement with the XRD results as a broad peak was observed. In contrast, the particles in the sol–gel combustion sample were sintered and fully formed a large crystalline (Fig. 2(D)). This can ascribe to the fact that the mesoporous SBA-15 would protect particle growth during the synthesis process by stabilizing and...
prohibiting agglomeration of CuFe$_2$O$_4$. However, Fig. 2(B) demonstrates the incomplete pore filling of SBA-15 by CuFe$_2$O$_4$. This result was in good agreement with the XRD pattern where the intensity of peaks could even vanish if pores were totally filled. Thus, the bright spots represented the pores of SBA-15, which were not filled with the CuFe$_2$O$_4$ particles. In addition, the incomplete pore filling is due to the faster ramping rate of calcination temperature that led to building up pressure from the decomposotion of precursor, hence, the capillary flow of the melting precursor salts into the mesoporous of SBA-15 silica template was restrained.$^{10,11}$

### 3.1.3 BET surface area analysis of CuFe$_2$O$_4$ spinel.

The textural properties of materials were performed using N$_2$ sorption isotherm and displayed in Fig. 3. The BET surface area of the synthesized catalysts is tabulated in Table 1. From the result, the N$_2$ sorption isotherms of CuFe$_2$O$_4$/SBA-15 composite displays a typical type IV adsorption according to IUPAC classification. A capillary condensation at a relative pressure of 0.5–0.9 was attributed to hysteresis loop type H1, evidently the mesoporosity of materials compared to SBA-15.$^{23}$ The CuFe$_2$O$_4$/SBA-15 composite possessed a lower surface area than SBA-15, which is probably due to partial filling of mesopores during the generation of spinel oxide as illustrated in the TEM images. In case of spinel oxide samples, the isotherms of CuFe$_2$O$_4$-TP and CuFe$_2$O$_4$-SG were significantly different from CuFe$_2$O$_4$/SBA-15 composite due to drastic reduction of the surface area. There is, however, a small hysteresis loop with isotherm being shifted to the high relative pressure for CuFe$_2$O$_4$-TP confirmed a mesoporous type of material. This could suggest that some parts of the space occupied by siliceous SBA-15 were removed and a broad space between needle-like spinel oxide was created, resulting in its higher surface area when compared with CuFe$_2$O$_4$ synthesized by sol-gel combustion.$^{8}$ It could be claimed that the template method was favourable for maintaining the nanocrystallite size with porosity even when the template was removed.

#### 3.1.4 Magnetization of CuFe$_2$O$_4$ spinel.

The magnetic hysteresis loops of samples with different preparation methods are also investigated to evaluate finite size effects (the high surface-to-volume ratio, different crystal structures and magnetic nanoparticles) on their catalytic activity. The surface-to-volume ratio of the nanoparticles show more significant effect on chemical reactivation rate compared to that of the corresponding bulk material.$^{14}$ Normally, the surface magnetization depends on the size of the particle contributed by the surface uncompensated spins. From Fig. 4, the result revealed that only the CuFe$_2$O$_4$-SG sample exhibited the magnetic hysteresis loops, while the CuFe$_2$O$_4$-TP and CuFe$_2$O$_4$/SBA-15 samples showed superparamagnetic curves at room temperature.

![Fig. 3 N$_2$ adsorption–desorption isotherms of spinel and SBA-15 samples; adsorption (black) and desorption (white).](image)

**Table 1** BET and structural parameters obtained from EXAFS data for the three calcined CuFe$_2$O$_4$ catalysts$^{a}$

| Sample name     | Paths                  | Occupancy (%) | R (Å)  | $\sigma^2$ | BET* (m$^2$ g$^{-1}$) |
|-----------------|------------------------|---------------|--------|------------|-----------------------|
| CuFe$_2$O$_4$-SG| Fe-O$_{\text{octa}}$ 2.8| 48.2          | 1.96   | 0.006      | 1                     |
|                 | Fe-Fe$_{\text{octa}}$ 2.8| 92.9          | 2.09   | 0.009      |                       |
|                 | Fe-Cu$_{\text{octa}}$ 3.8| 37.7          | 3.38   | 0.003      |                       |
|                 | Fe-O$_{\text{octa}}$ 2.8| 93.8          | 3.82   | 0.009      |                       |
|                 | Fe-O$_{\text{tetra}}$ 1.2| 51.8          | 2.05   | 0.020      |                       |
|                 | Fe-Cu$_{\text{tetra}}$ 3.8| 36.3          | 3.63   | 0.004      |                       |
|                 | Fe-Cu$_{\text{tetra}}$ 3.8| 32.0          | 3.82   | 0.009      |                       |
|                 | Fe-O$_{\text{2tetra}}$ 6.5| 38.6          | 3.86   | 0.040      |                       |
| CuFe$_2$O$_4$-TP| Fe-O$_{\text{octa}}$ 3.8| 63.4          | 1.96   | 0.002      | 30                    |
|                 | Fe-Cu$_{\text{octa}}$ 3.8| 29.4          | 2.94   | 0.009      |                       |
|                 | Fe-Cu$_{\text{tetra}}$ 3.8| 41.1          | 3.41   | 0.004      |                       |
|                 | Fe-O$_{\text{tetra}}$ 3.8| 47.4          | 3.47   | 0.004      |                       |
|                 | Fe-Cu$_{\text{tetra}}$ 3.8| 36.6          | 1.84   | 0.002      |                       |
|                 | Fe-Cu$_{\text{tetra}}$ 2.2| 73.3          | 3.73   | 0.004      |                       |
|                 | Fe-O$_{\text{2tetra}}$ 2.2| 39.2          | 3.92   | 0.004      |                       |
|                 | Fe-O$_{\text{2tetra}}$ 4.4| 83.8          | 3.83   | 0.004      |                       |
| CuFe$_2$O$_4$/SBA-15| Fe-O$_{\text{octa}}$ 5.1| 84.3          | 1.96   | 0.014      | 336                   |
|                 | Fe-Cu$_{\text{octa}}$ 5.1| 29.8          | 2.98   | 0.001      |                       |
|                 | Fe-Cu$_{\text{tetra}}$ 5.1| 39.9          | 3.39   | 0.004      |                       |
|                 | Fe-O$_{\text{2tetra}}$ 5.1| 44.4          | 3.44   | 0.027      |                       |
|                 | Fe-O$_{\text{2tetra}}$ 0.6| 15.7          | 1.96   | 0.001      |                       |
|                 | Fe-Cu$_{\text{tetra}}$ 0.9| 63.6          | 3.64   | 0.002      |                       |
|                 | Fe-Cu$_{\text{tetra}}$ 0.9| 52.0          | 3.52   | 0.011      |                       |
|                 | Fe-O$_{\text{2tetra}}$ 1.9| 63.9          | 3.69   | 0.002      |                       |

*Where $N$ = coordination number, $R$ = distance to neighboring atom, $^*$ SBA-15 = 685 m$^2$ g$^{-1}$. 

![Fig. 4 Characteristic hysteresis curves of magnetization (M) versus applied magnetic field (H) for calcined CuFe$_2$O$_4$/SBA-15, CuFe$_2$O$_4$-TP, and CuFe$_2$O$_4$-SG.](image)
temperature. This result agrees well with the results reported by Laokul et al.\textsuperscript{25} and Yadav et al.\textsuperscript{26} According to XRD and TEM results, the particle sizes of the CuFe\textsubscript{2}O\textsubscript{4} species in CuFe\textsubscript{2}O\textsubscript{4-TP} and CuFe\textsubscript{2}O\textsubscript{4}/SBA-15 were very small, only observing in a few nanometres. It is obvious that these nanoparticles reached superparamagnetic sizes. The magnetizations of samples were in the order of CuFe\textsubscript{2}O\textsubscript{4-SG} > CuFe\textsubscript{2}O\textsubscript{4-TP} > CuFe\textsubscript{2}O\textsubscript{4}/SBA-15. This could possibly result from the effect of particle size. As particle size decreases, the surface-to-volume ratio increases. The noncollinearity of the spins from the surface of the nanoparticles is reflected in the decrease of the saturation magnetization, as compared to that of the bulk material. The decrease of the saturation magnetization can be higher in ferrimagnetic nanoparticles, where the exchange interaction takes place over a certain distance. However, this effect is less pronounced when the nanoparticles are very small, due to the dominance of the superparamagnetic sizes. The magnetizations of samples were almost equally distributed between the octahedral and tetrahedral sites. For CuFe\textsubscript{2}O\textsubscript{4-TP} and CuFe\textsubscript{2}O\textsubscript{4}/SBA-15 samples, 63.4% and 84.3% of Fe ions respectively occupied octahedral sites, while Cu ions were partly located at the octahedral sites. It is clear that the cation distribution in CuFe\textsubscript{2}O\textsubscript{4} spinel nanoparticles was strongly affected by preparation methods.

### 3.1.5 Local structure and cation distribution of CuFe\textsubscript{2}O\textsubscript{4} spinel

The Fe K-edge XANES spectra of the reference compounds and the calcined CuFe\textsubscript{2}O\textsubscript{4} catalysts are presented in Fig. 5. The XANES feature at Fe K-edge of standard CuFe\textsubscript{2}O\textsubscript{4} sample was similar to that of Fe\textsubscript{3}O\textsubscript{4} with the higher edge threshold energy. This indicated that the CuFe\textsubscript{2}O\textsubscript{4} sample contained higher amount of Fe\textsuperscript{3+} species. However, as observed from all samples, there was no significant difference in spectral features compared to CuFe\textsubscript{2}O\textsubscript{4} standard materials. Small absorption peaks at the pre-edge region were observed due to the electric quadrupole-allowing transitions of Fe species confirming the existence of Fe ions in the tetrahedral and octahedral sites. Normally, cuprospinel is an inverse spinel where Cu\textsuperscript{2+} only occupies in octahedral sites and iron ions (Fe\textsuperscript{2+} and Fe\textsuperscript{3+}) are equally distributed between the octahedral and tetrahedral sites in the structure.\textsuperscript{23} The Fe\textsuperscript{2+} ions prefer to occupy octahedral sites while the Fe\textsuperscript{3+} ions prefer to occupy tetrahedral sites.\textsuperscript{24}

In EXAFS spectra (Fig. 6), the intensity of the first peak arises from the scattering of the nearest neighbour oxygen anions (Fe–O) while the second large peak is caused by the scattering of the Fe and Cu, possibly located in both tetrahedral and octahedral sites. The interatomic distance of the Fe–O and Fe–Fe bonds was insignificantly different in each sample. However, a variation of relative peak heights of these two major peaks was observed, which would indicate a significant change in the local structural environment of Fe ions in the CuFe\textsubscript{2}O\textsubscript{4} catalysts prepared with three different synthesis methods. From the EXAFS analysis, the quantitative information on the cation distribution was illustrated in Table 1. In CuFe\textsubscript{2}O\textsubscript{4-SG}, Fe ions were almost equally distributed between the octahedral and tetrahedral sites, and all Cu ions should be located at the octahedral sites. For CuFe\textsubscript{2}O\textsubscript{4-TP} and CuFe\textsubscript{2}O\textsubscript{4}/SBA-15 samples, 63.4% and 84.3% of Fe ions respectively occupied octahedral sites, which would suggest that Cu ions were partly located at the octahedral and tetrahedral sites. It is clear that the cation distribution in CuFe\textsubscript{2}O\textsubscript{4} spinel nanoparticles was strongly affected by preparation methods.

![Fig. 5](image1)

**Fig. 5** Fe K-edge XANES spectra of CuFe\textsubscript{2}O\textsubscript{4-SG}, CuFe\textsubscript{2}O\textsubscript{4-TP} and CuFe\textsubscript{2}O\textsubscript{4}/SBA-15 samples compared with standard materials of CuFe\textsubscript{2}O\textsubscript{4}, Fe\textsubscript{3}O\textsubscript{4}, Fe\textsubscript{2}O\textsubscript{3}, CuFeO\textsubscript{2}, and FeO.

![Fig. 6](image2)

**Fig. 6** (A) EXAFS oscillations, $k^2 \chi (k)$, and (B) their Fourier transforms of CuFe\textsubscript{2}O\textsubscript{4}/SBA-15, CuFe\textsubscript{2}O\textsubscript{4-TP} and CuFe\textsubscript{2}O\textsubscript{4-SG}.
3.1.6 Reduction behaviours of CuFe₂O₄ spinel. In order to understand the transformation of CuFe₂O₄ spinel under the reduction condition, the H₂-TPR was particularly employed for the study. Illustrated in Fig. 7, two pronounced reduction regions were observed for each sample. It has been reported that the reduction peak at lower temperature range of 180–420 °C was responsible for the reduction of CuFe₂O₄ to Cu⁰ and Fe₃O₄ species. Subsequently, Fe₃O₄ is reduced to Fe⁰ particles at higher temperature (420–750 °C). These results were in close agreement with those reported in literatures. According to these results, it was difficult to identify a specific species during these two reduction steps. Normally, CuFe₂O₄ was mainly reduced to Cu and FeO₃ by removing of the lattice oxygen under H₂ atmosphere, then FeO₃ was rapidly reduced to Fe₂O₃. For comparison with individual TPR profiles, it was found that CuFe₂O₄_TP and CuFe₂O₄_SG showed almost the same reduction temperature compared to CuFe₂O₄/SBA-15. The unsupported CuFe₂O₄ required higher temperature for its reduction process. In contrast, CuFe₂O₄/SBA-15 composite has a lowest reduction temperature for reducing CuFe₂O₄ to metallic copper and Fe₂O₃ oxide, indicating that metal supported materials was easier reduced than the bulk (unsupported) metal oxides. This phenomenon can be explained in term of specific surface area and H₂ spillover. When the SBA-15 surface was completely covered by the dissociative adsorbed hydrogen atom, it spilled over the surface of SBA-15 and migrated to react with CuFe₂O₄. Hence, it increased the reduction to a higher rate. The lower reduction temperature of CuFe₂O₄/SBA-15 might be a result of the generation of nanometallic Cu active sites, which enhanced the catalytic properties.

To further investigate the reductive species in the reduced samples, Cu and Fe K-edges XANES spectra were collected and have been displayed in Fig. 8. In the Cu K-edge, the edge energies of the reduced samples are the same as the Cu foil, indicating the presence of metallic copper. A closer look at XANES spectra, the white line feature and shoulder were observed in CuFe₂O₄_SG, indicating dominating copper oxides in the samples. Here, the intense white line peak and the dominant shoulder in the CuFe₂O₄_SG could be assigned to the 1s → 4p transition of Cu₂O. To analyze the composition of copper species in the samples, the linear combination fit analysis was applied. The quantification by linear combination fitting yields a mixture of metallic Cu, Cu₂O and CuO in each sample (Fig. 8). It was found that the copper ions in CuFe₂O₄/SBA-15 were highly reduced to the state of zero, while the CuFe₂O₄_SG was partially reduced at 350 °C. In case of Fe K-edge, the XANES fit of Fe K-edge revealed that the catalysts were fully remained in the oxide forms. The structure of iron was gradually change from CuFe₂O₄ to typically iron oxides like FeO, Fe₂O₃, and Fe₃O₄. These results are consistent with the H₂-TPR which ascribe that iron species in CuFe₂O₄ spinel are more resistant toward reduction.

Fig. 7 TPR profiles of (A) CuFe₂O₄_SG, (B) CuFe₂O₄_TP, and (C) CuFe₂O₄/SBA-15.

Fig. 8 Linear combination fitted-XANES spectra of CuFe₂O₄/SBA-15, CuFe₂O₄_TP, and CuFe₂O₄_SG after reduction under H₂ atmosphere.
3.2 Testing of CuFe₂O₄ spinel for steam reforming of methanol (SRM)

Prior to demonstrate the catalytic performance, the blank test of pure siliceous SBA-15 was carried out with the same space velocity. The result showed a negligible catalytic activity of the siliceous SBA-15. In Fig. 9, the catalytic results of each catalyst were represented in the term of H₂ production rate and methanol conversion as a function of temperature. After H₂ reduction at 350 °C, it was observed that the rate of methanol conversion and H₂ production rate per gram of catalyst were in order of CuFe₂O₄_SG > CuFe₂O₄/SBA-15 > CuFe₂O₄_TP. However, when considering the H₂ production rate per gram of active CuFe₂O₄ spinel, the CuFe₂O₄/SBA-15 catalyst showed the highest H₂ production rate (~5500 mol kg⁻¹ CuFe₂O₄ h⁻¹) at the reaction temperature of 375 °C. It is suggested that the improved turnover rate of CuFe₂O₄/SBA-15 was due to the high dispersion of copper spinel particles on the SBA-15 support. Moreover, the mesoporous SBA-15 silica support could obstruct the sintering process of nano-metallic copper during the reaction and enhanced the catalytic reaction.

Comparison between CuFe₂O₄_SG and CuFe₂O₄_TP catalysts, it was very interesting that the lower surface of CuFe₂O₄_SG provided the highest catalytic activity (100% conversion and H₂ production rate of ~1600 mol kg⁻¹ Cat h⁻¹ at 375 °C) compared to the higher surface area of CuFe₂O₄_TP. The CuFe₂O₄_TP exhibits the lowest yield of H₂ production rate (<10 mol kg⁻¹ Cat h⁻¹) with % methanol conversion of less than 1%. This result suggested that almost no metallic copper was generated from the reduction of the needle-like CuFe₂O₄_TP. To verify this concept, the catalytic evaluation of CuFe₂O₄_SG and CuFe₂O₄_TP was also studied under the reduction temperature of 420 °C as indicated by TPR results that CuFe₂O₄_SG and CuFe₂O₄_TP catalysts could be completely reduced at temperature higher than 350 °C. However, the catalytic activity of these two catalysts after reducing at 420 °C was dramatically decreased in their performance. This could be attributed to the fact that the reduced copper was sintered to form bulky particles during the reduction process. Hence, the reduction temperature at 350 °C was enough for reduction of CuFe₂O₄_SG and CuFe₂O₄_TP in this reaction. Nevertheless, it was an unexpected result that the needle-like CuFe₂O₄_TP was not active for the methanol steam reforming. This issue should be further investigated and reported in next.

4. Conclusions

The preparation methods strongly affected on cation distribution and physicochemical properties of CuFe₂O₄. The CuFe₂O₄ supported on SBA-15 exhibited the superparamagnetic behaviour with well-crystallized tetragonal structure and high reducibility. However, upon removal of SBA-15 from the CuFe₂O₄/SBA-15, the needle-like of CuFe₂O₄_TP with high surface area, but the reduction requires a high temperature as well as sol–gel combustion method.

Conflicts of interest

There are no conflicts to declare.

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