

High hydrothermal stability of Cu–SAPO-34 catalysts for the NH₃-SCR of NO_x

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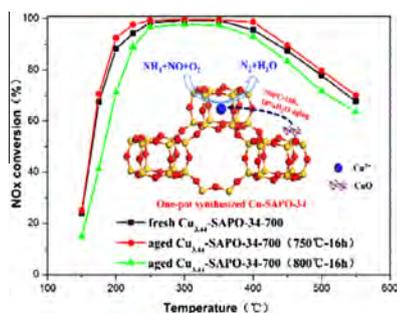
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HIGHLIGHTS

- Relatively high calcination temperature was favorable for the deNO_x catalytic activity of Cu–SAPO-34.
- Hydrothermal treatment increased NH₃-SCR activity due to the migration of CuO to isolated Cu²⁺ as well as the decreased activation barrier.
- High hydrothermal stability depend on the maintenance of active Cu²⁺, crystallinity and the acidity after harsh treatment.

GRAPHICAL ABSTRACT

Cu–SAPO-34 catalysts prepared by the one-pot hydrothermal synthesis method showed high activity and outstanding hydrothermal stability due to the maintenance of isolated Cu²⁺, the crystallinity and acidity as well as low activation barrier.



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ABSTRACT

The effects of calcination temperature, Cu source and co-template contents on the activity and hydrothermal stability of Cu–SAPO-34 catalysts prepared by the one-pot hydrothermal synthesis method were investigated. Appropriate calcination temperature was important for the deNO_x activity of Cu–SAPO-34 catalyst. The amount of Cu source and co-template in the resulting gel could impact the Cu loading and the crystallization of the final Cu–SAPO-34. Preferably, hydrothermal treatment of the Cu–SAPO-34 catalyst with Cu loading ca. 3.44% at 750 °C for 16 h increased the NH₃-SCR activity slightly due to migration of CuO to isolated Cu²⁺ and the maintenance of the crystallinity as well as the acidity. Moreover, the catalyst aged at 800 °C for 16 h maintained >90% NO_x conversion from 225 to 400 °C, and the crystallinity, active Cu²⁺ species and acidity in Cu–SAPO-34 were not destroyed completely after such harsh treatment. Meanwhile, kinetic analysis over fresh and aged Cu–SAPO-34 samples were conducted and the conclusion were in agreement with the NH₃-SCR tests, H₂-TPR and NH₃-TPD results.

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1. Introduction

Pollutants, especially NO_x from diesel engine exhaust, are the main sources of acid rain, haze and photochemical smog. Reducing NO_x emissions is important for the improvement of air quality. The selective catalytic reduction of NO_x by NH₃ (NH₃-SCR) is one of the most promising technologies for NO_x emission control for diesel

engines [1]. Due to the unstable operating conditions of diesel engines and periodically increasing the exhaust temperature to replicate diesel particulate filter (DPF) regeneration, excellent catalysts should meet the following essential requirements: good SCR activity over a wide operating temperature window, high selectivity for N_2 , and outstanding hydrothermal stability.

Copper-based zeolites have been suggested as efficient catalysts for the NH_3 -SCR reaction [2–5], and Cu-ZSM-5 was proposed to be the most promising [6,7]. However, the structure of the Cu-ZSM-5 catalyst collapsed easily after high temperature hydrothermal aging, leading to a conspicuous decline in NH_3 -SCR activity [8]. Recently, Cu-CHA catalysts, (Cu supported on microporous zeolites (~0.38 nm pore size)), such as Cu-SSZ-13 and Cu-SAPO-34, have attracted much attention due to their higher NH_3 -SCR activity and better hydrothermal stability compared to large or mesopore Cu-based zeolite catalysts [9,10].

Generally, Cu-CHA catalysts are prepared by ion-exchange methods (including aqueous solution and solid-state ion-exchange). The aqueous solution ion exchange method involves a two-step exchange process, filtering, washing and calcining, greatly limiting its application [11,12]. The solid-state ion exchange (SSIE) method is carried out under rather harsh conditions (calcining at 800 °C), leading to decomposition of the framework [13,14]. Recently, one-pot hydrothermal synthesis method used low-cost copper-tetraethylenepentamine (Cu-TEPA) as Cu source and template to prepare Cu-CHA directly which was much simpler to perform, and might be applicable at industrial scale [15]. Xie et al. [16] reported that the NO_x conversion of one-pot synthesized $Cu_{3.8}$ -SSZ-13 reached ca. 85% at 200 °C and maintained a level above 90% until 550 °C. Raquel et al. [17,18] reported that the NO conversion of $Cu_{3.3}$ -SAPO-34 sample maintained a level above 90% between 200 and 450 °C and the NO conversion decreased about 10% over the whole temperature range after hydrothermal aging at 750 °C for 13 h. However, it is still a blank to investigate the one-pot hydrothermal synthesized Cu-SAPO-34 catalysts under more severe hydrothermal deactivation conditions. For instance, hydrothermal aging at 800 °C for 16 h can compare with a 135,000 mile vehicle-aged SCR catalyst [19].

In this study, Cu-SAPO-34 catalysts were prepared using Cu-TEPA as Cu source and propylamine (PA) as co-template. Three key preparation parameters, including calcination temperature, Cu source and co-template amounts, were optimized systematically, and Cu-SAPO-34 catalysts with high NO_x removal efficiency, N_2 selectivity and hydrothermal stability were synthesized successfully. By using N_2 adsorption/desorption, XRD, H_2 -TPR and NH_3 -TPD, the origin of the high activity and excellent hydrothermal stability was investigated in detail.

2. Experimental

2.1. Catalyst synthesis

A series of Cu-SAPO-34 catalysts was prepared using Cu-TEPA as Cu source, PA as co-template, pseudoboehmite as the Al source, 85% phosphoric acid as the P source, and fumed silica as the Si source. The molar composition of the synthesized gel was: 1Al:1P:0.25Si:(0.03–0.20)Cu-TEPA:(1.2–1.8)PA:40 H_2O . The resulting gel was transferred to an autoclave with a Teflon liner, and crystallized for 72 h at 180 °C. The crystalline products were filtered and washed with distilled water, and dried at 100 °C overnight.

In order to investigate the effect of calcination temperature, an initial Cu-SAPO-34 sample with Cu loading of about 5% was prepared. The catalysts were then calcined in air for 5 h to remove the organic templates at temperatures of 500, 600, 700 or 800 °C, and designated as Cu_x -SAPO-34-T, where T represented the calcination temperature.

By adjusting the content of Cu-TEPA with constant PA/Al = 1.6, Cu-SAPO-34 samples with different Cu loadings can be obtained (Table 1). By changing the input of PA, with constant Cu-TEPA/Al = 0.06, Cu-SAPO-34 samples with varying properties can also be obtained (Table 2). These initial powder samples were calcined in air at 700 °C, and designated as Cu_x -SAPO-34 with x representing the Cu loading.

To investigate the hydrothermal stability of Cu-SAPO-34 samples, the catalysts were hydrothermally aged at 750 °C and 800 °C for 16 h in a flow of 10% H_2O /air.

2.2. NH_3 -SCR activity measurement

SCR activity tests of the sieved powder catalysts were carried out in a fixed-bed quartz flow reactor at atmospheric pressure. The reaction conditions were controlled as follows: 500 ppm NO , 500 ppm NH_3 , 5 vol.% O_2 , balance N_2 and 500 mL/min total flow rate. During the performance tests, about 60 mg catalyst was used, yielding a rather high GHSV of 400,000 h^{-1} . The effluent gas, including NO , NH_3 , NO_2 , and N_2O , was continuously analyzed by an online NEXUS 670-FTIR spectrometer equipped with a heated, low volume (0.2 L) multiple-path gas cell (2 m). The FTIR spectra were collected throughout and the results were recorded when the SCR reaction reached a steady state. In this study, we defined the N_2 selectivity as the percentage of N atoms in the feed gas $NO + NH_3$ not changing into N_2O and NO_2 in the SCR reaction process. Then, the NO_x conversion and N_2 selectivity were calculated according to the following equations:

$$NO_x \text{ conversion} = \left(1 - \frac{[NO]_{out} + [NO_2]_{out}}{[NO]_{in} + [NO_2]_{in}} \right) \times 100\%$$

$$N_2 \text{ selectivity} = \frac{[NO]_{in} + [NH_3]_{in} - [NO_2]_{out} - 2[N_2O]_{out}}{[NO]_{in} + [NH_3]_{in}} \times 100\%$$

2.3. Characterization

N_2 adsorption/desorption isotherms of the Cu-SAPO-34 samples were measured at –196 °C using a Quantachrome Quadrasorb SI-MP to obtain the specific surface area and pore volume. Prior to the N_2 physical adsorption, the samples were degassed at 300 °C for 5 h. Surface areas and pore volumes were determined by the t-plot method.

Powder X-ray diffraction (XRD) measurements were carried out on a computerized PANalytical X'Pert Pro diffractometer with $Cu K\alpha$ ($\lambda = 0.15406$ nm) radiation. The data of 2θ from 5° to 40° were collected with the step size of 0.02°.

Temperature-programmed reduction with hydrogen (H_2 -TPR) experiments were carried out on a Micromeritics AutoChem 2920 chemisorption analyzer. The interference of H_2O was eliminated by using a liquid nitrogen cold trap before the detector. 50 mg of the samples in a quartz reactor were pre-treated in air with the flow rate of 50 mL/min at 500 °C for 1 h. After the catalyst was cooled down to room temperature, H_2 -TPR was performed in 10 vol.% H_2 /Ar gas flow of 50 mL/min at a heating rate of 10 °C/min to 900 °C.

Table 1

Cu loadings of the Cu_x -SAPO-34 by adjusting Cu-TEPA contents in the resulting gel.

Samples	Cu-TEPA/Al	Cu, wt.%
$Cu_{2.71}$ -SAPO-34	0.03	2.71
$Cu_{3.44}$ -SAPO-34	0.06	3.44
$Cu_{6.62}$ -SAPO-34	0.15	6.62
$Cu_{8.84}$ -SAPO-34	0.20	8.84

Table 2
Properties of Cu_x-SAPO-34 samples prepared with different PA inputs in the resulting gel.

Samples	PA/Al	pH of resulting gel	Cu, wt.%	Crystalline phases
Cu _{2.96} -SAPO-34	1.2	6.0	2.96	Amorphous + SAPO-34
Cu _{3.20} -SAPO-34	1.4	7.0	3.20	Amorphous + SAPO-34
Cu _{3.44} -SAPO-34	1.6	8.0	3.44	SAPO-34
Cu _{5.15} -SAPO-34	1.8	9.5	5.15	SAPO-34

Ultraviolet visible diffuse reflectance spectrum (UV-vis-DRS) were recorded at room temperature using a Hitachi UV-3010 spectrometer equipped with a diffuse reflectance accessory in the 200–800 nm wavelength range. BaSO₄ was used as the reference sample.

X-ray photoelectron spectroscopy (XPS) with Al Kα radiation (1486.7 eV) was used to analyze the atomic state of copper species on the catalysts surface (Axis Ultra, Kratos Analytical Ltd). The C 1s peak at 284.8 eV was used as an internal standard for peak position measurement.

Temperature-programmed desorption with ammonia (NH₃-TPD) experiments were performed in the same instrument as H₂-TPR, equipped with a quadrupole mass spectrometer (MKS Cirrus) to record the signals of NH₃ (m/z = 17 for NH₃). The interference of H₂O was eliminated by using a cold trap before the detector. Prior to TPD experiments, 50 mg of the samples were pre-treated in air at a flow rate of 50 mL/min at 500 °C for 1 h, and then cooled down to room temperature (30 °C). The samples were then exposed to a flow of 2500 ppm NH₃/Ar (50 mL/min) at 30 °C for 1 h, followed by Ar purging for 1 h. Finally, the temperature was raised to 900 °C in Ar at the rate of 10 °C/min.

2.4. Kinetic studies

NH₃-SCR kinetic tests over fresh and aged catalysts were evaluated in a differential reactor with 4 mm inner diameter at atmospheric pressure. About 10 mg of the catalyst diluted with 40 mg of quartz sand was loaded with a catalyst bed length of ca. 2 mm. Under the different conditions, the NO_x conversion was kept less than 20% in the temperature range tested. To eliminate the effects of diffusion, a total gas flow rate of 500 mL/min and particle size of 40–60 mesh were used. The gas mixture was composed of 500 ppm NO, 500 ppm NH₃, 5 vol.% O₂, and the balance N₂. The effluent gas was also analyzed by an online NEXUS 670-FTIR spectrometer as described above. NO_x reduction rates (mol/g/s) were calculated as:

$$r_{\text{NO}_x} = X_{\text{NO}_x} Y_{\text{NO}_x, \text{in}} V_{\text{gas}} / m_{\text{cat}}$$

where X_{NO_x} is the NO_x conversion, $Y_{\text{NO}_x, \text{in}}$ is the NO_x molar fraction in the inlet gas, V_{gas} is the total flow rate in moles per second, and m_{cat} is the mass of the catalyst in grams.

3. Results and discussion

3.1. Effect of calcination temperature

3.1.1. NH₃-SCR activity and N₂ selectivity

The NO_x conversion and N₂ selectivity of Cu₅-SAPO-34-T catalysts with different calcination temperatures are shown in Fig. 1. It can be observed that calcination temperature had a remarkable influence on the catalytic performance of Cu₅-SAPO-34 catalyst. Among the catalysts calcined at different temperatures, Cu₅-SAPO-34-700 exhibited superior low-temperature SCR activity and a broad temperature window for NO_x removal, with almost 100% NO_x conversion in the range of 250–400 °C. The steady state NO_x conversion below 350 °C increased with increasing calcination

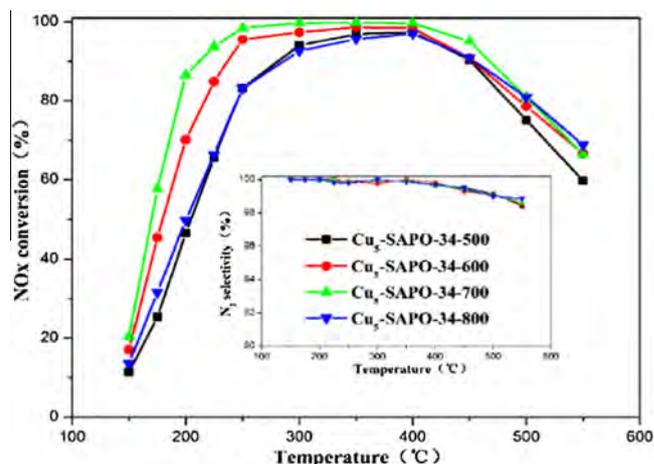


Fig. 1. NO_x conversion and N₂ selectivity over Cu₅-SAPO-34-T samples with varied calcination temperatures.

temperature from 500 to 700 °C, and then decreased for higher calcination temperature at 800 °C. There was little effect on NH₃-SCR activity above 350 °C. The N₂ selectivity of all Cu₅-SAPO-34-T samples were higher than 98% in the whole reaction temperature range. These results suggested clearly that appropriate calcination temperature was important for the catalytic activity of Cu-SAPO-34. Gao et al. [13] studied solid-state reaction between SAPO-34 and CuO under different calcination temperatures and times and found that a portion of CuO (~67%) converted to isolated Cu²⁺ ions after the SSIE treatment at 800 °C for 16 h. It suggested that the copper species in Cu-SAPO-34 can be redistributed to yield an active catalyst upon high temperature treatment. However, copper species distribution of Cu₅-SAPO-34-T samples did not change distinctly (Fig. S1), suggesting that the deNO_x catalytic activity differences among Cu₅-SAPO-34-T samples mainly attributed to the varying structural properties.

Furthermore, the NO_x conversion over hydrothermally aged Cu₅-SAPO-34-T catalysts was investigated. As shown in Fig. 2(c), the deNO_x catalytic activity of aged Cu₅-SAPO-34-700 (750 °C–16 h) did not significantly change at low temperatures and slightly decreased at high temperatures. By comparing with all the fresh and aged samples, it was seen clearly that the Cu₅-SAPO-34-700 showed excellent SCR activity and hydrothermal stability, while the Cu₅-SAPO-34-800 performed inferiorly especially under harsh treatment.

3.1.2. N₂ adsorption/desorption

Table 3 shows the specific surface area and pore volume of the Cu₅-SAPO-34-T samples calcined at different temperatures. The specific surface area of the Cu-SAPO-34 catalysts were strongly affected by the calcination temperatures and increased monotonically from 501.1 m²/g to 566.6 m²/g with the calcination temperature rising from 500 to 700 °C. When catalysts were calcined at higher temperatures (800 °C), the surface area decreased. It is widely accepted that the surface area of zeolites decreases with the increase of the calcination temperatures, as observed for Fe-

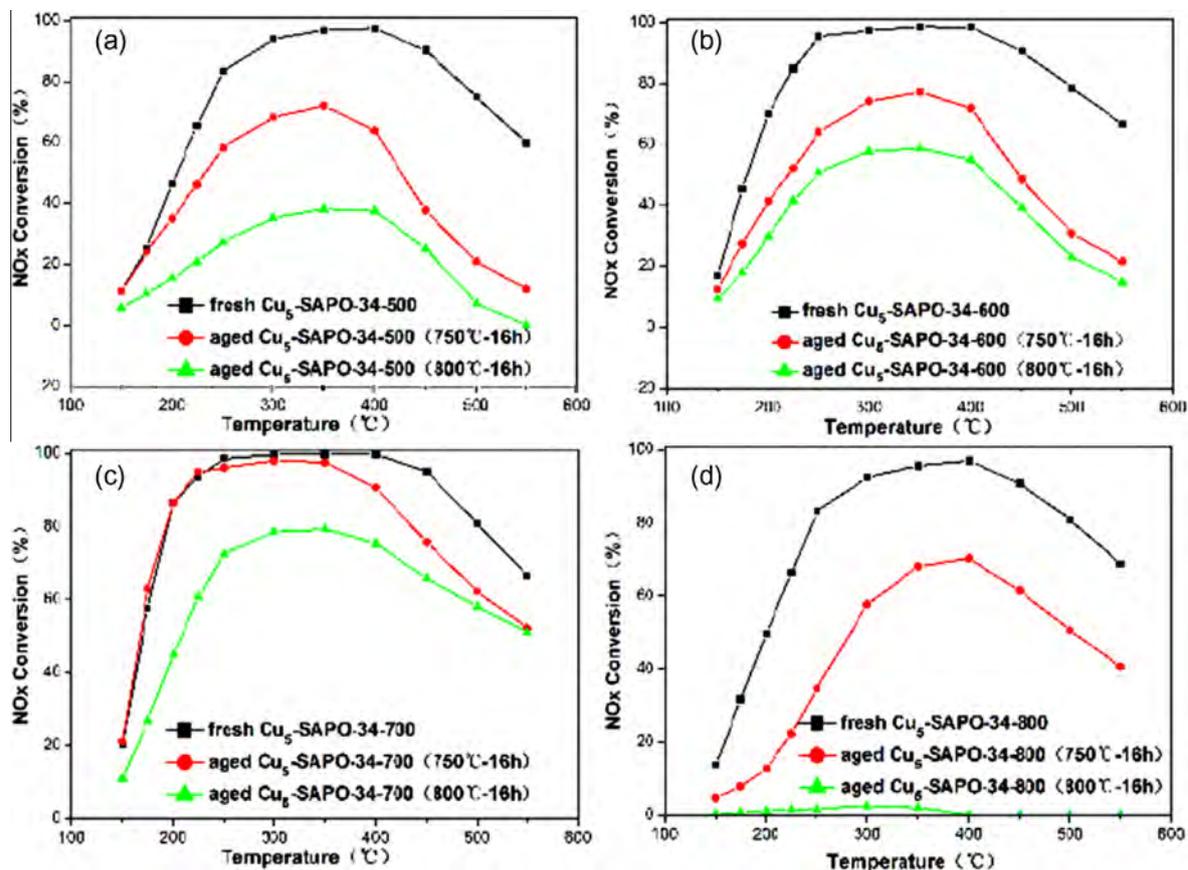


Fig. 2. NOx conversion over fresh and aged Cu₅-SAPO-34-T samples with varied calcination temperatures.

Table 3

Properties of the Cu₅-SAPO-34-T samples calcined at different temperatures.

Samples	Specific surface area, m ² /g	Pore volume, cm ³ /g
Cu ₅ -SAPO-34-500	501.1	0.26
Cu ₅ -SAPO-34-600	506.8	0.26
Cu ₅ -SAPO-34-700	566.6	0.29
Cu ₅ -SAPO-34-800	480.7	0.25

HBEA and Mo/HZSM-5 [20,21]. However, the Cu-SAPO-34 catalyst in the present study differed from those zeolites possibly due to the pore structure forming gradually during the calcination process and reached a maximum. Because of the same Cu loading and similar copper species distribution, the activity of Cu₅-SAPO-34-T eliminated the effect of Cu species which could influence the SCR activity. As shown in Fig. 1, the Cu₅-SAPO-34-700 sample with the largest surface area and pore volume, which can supply the largest inner surface area for the SCR reaction, showed the highest NOx conversion in the SCR reaction. Li et al. [22] also indicated that the excellent catalytic activity was attributed to the large specific surface area.

3.1.3. Powder XRD

To further confirm the peak intensity changes and the integrity of the CHA zeolite structure under different calcination temperatures, the XRD patterns of Cu₅-SAPO-34-T samples were obtained and are shown in Fig. 3. Typical CHA structure diffraction peaks (see <http://www.iza-structure.org/databases> for standard XRD patterns) were observed in all samples. Gao et al. [13] reported that a weak diffraction feature of SiO₂ appeared for the sample aged at 800 °C for 1, 5 and 16 h, suggesting some decomposition. Never-

theless, there was no evident that CHA structure decomposed for sample calcined at 800 °C for 5 h in our work, indicating that the physical structure of Cu₅-SAPO-34-T sample was stable under the current calcination temperatures.

Increase in intensity of the XRD peaks represented the increase in the crystallinity [23]. The crystallinity increased with increasing calcination temperature from 500 to 700 °C, and then declined when the calcination temperature was 800 °C, which conformed well with the trend of SCR activity. Ye et al. [24] reported that the Ag-SAPO-34 which had high degree of crystallinity, showed good catalytic activity. The Cu₅-SAPO-34 catalyst with the highest degree of crystallization, surface area and pore volume could be obtained by calcination at an optimum temperature of 700 °C. Hence, the calcination temperature was chosen as 700 °C for further investigations in the present study.

3.2. Effects of Cu source and co-temple

3.2.1. NH₃-SCR activity and N₂ selectivity

SCR performance tests were conducted on samples with different Cu source contents in the resulting gel are shown in Fig. 4(a). The Cu_{2.71}-SAPO-34 sample containing the lowest Cu loading showed moderate activity, indicating that more active copper species were needed. When the Cu loading was 3.44%, the NOx conversion increased to 90% at 200 °C and maintained that level until 400 °C with the N₂ selectivity of more than 98% in the whole temperature range. With further increase of the Cu loading, the activity levelled off below 350 °C. However, at high temperatures (350–550 °C), the NOx conversion decreased rapidly when the Cu loading was above 3.44%, and a dramatic decline of NOx conversion and N₂ selectivity was observed over the Cu_{8.84}-SAPO-34 sample.

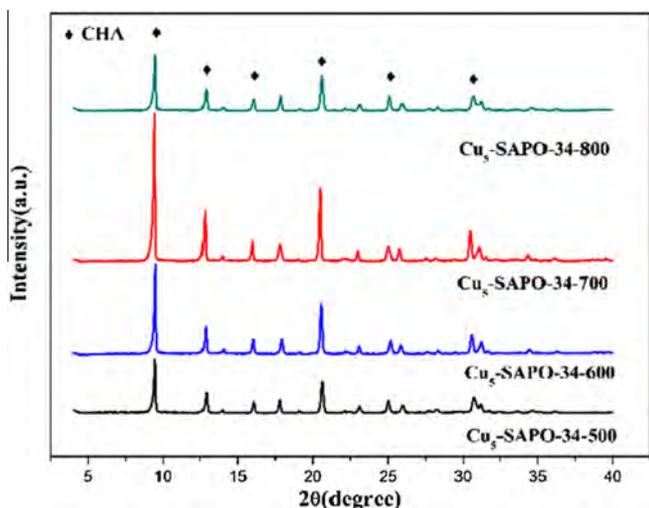


Fig. 3. Powder XRD patterns of $\text{Cu}_x\text{-SAPO-34-T}$ samples with varied calcination temperatures.

Most researchers consider that the active sites for $\text{NH}_3\text{-SCR}$ reactions are likely to be isolated Cu^{2+} ions at exchange sites [11,16,25,26] and that the CuO species could lead to non-selective oxidation of NH_3 and decreased $\text{NH}_3\text{-SCR}$ activity at high temperatures [14,27,28]. Table 4 shows that the amount of Cu^{2+} ions increased from 1.35% to 4.69% along with the improvement of the SCR activity below 350 °C. On the other hand, the amount of CuO increased from 0.35% to 2.07% along with the decrease of the SCR activity above 350 °C. With Cu loading from 3.44% to 6.62%, the NO_x conversion did not obviously improve, indicating that active sites were almost saturated on the $\text{Cu}_{3.44}\text{-SAPO-34}$ sample. The result showed that, there existed the best Cu loading rather than the Cu loading was the higher the better.

Properties of $\text{Cu}_x\text{-SAPO-34}$ samples prepared with different PA inputs in the resulting gel are shown in Table 2. With the increase of PA/Al ratio from 1.2 to 1.8, the pH of the gel increased from 6.0 to 9.5 with Cu loading increasing from 2.96% to 5.15%. The template agent such as PA used to fill the CHA framework and direct the synthesis of Cu-SAPO-34 , is a kind of strong alkalogenic organic amine. Therefore, the input of PA greatly influenced the chemical environment such as pH value of the resulting gel which could affect the degree of solubility and polymerization of raw materials and further impact the crystalline phase of the zeolite.

White et al. [29] indicated that zeolite synthesis is the process of inorganic oxide polymerization and deprotonation which depend greatly on the pH value using density functional theory (DFT). Lower pH value led to decreased solubility of amorphous silica in the silicate dimerisation reactions. As a result of shortage of the nuclear material in the solution at low pH, amorphous phase was obtained. Hence, Cu-SAPO-34 catalysts were only achieved at appropriate pH value.

As is shown in Fig. 4(b), with the increasing of PA inputs, the deNO_x catalytic activity of $\text{Cu}_x\text{-SAPO-34}$ samples improved below 350 °C and showed decline trend as a whole at high temperatures. Although the $\text{Cu}_{5.15}\text{-SAPO-34}$ (pH = 9.5) sample performed better than the $\text{Cu}_{3.44}\text{-SAPO-34}$ (pH = 8.0) sample below 350 °C, it cannot be concluded that both Cu-SAPO-34 samples were promising candidates for $\text{NH}_3\text{-SCR}$, since an excellent catalyst should possess high hydrothermal stability. Hence, hydrothermal treatments were conducted on the two samples to choose the better candidate, and further, the optimum synthesis condition, as shown in Section 3.3.

3.2.2. Powder XRD

The crystal structures of $\text{Cu}_x\text{-SAPO-34}$ samples were verified via XRD measurements. As shown in Fig. 5, $\text{Cu}_x\text{-SAPO-34}$ maintained the CHA framework structure basically after high temperature calcination. However, two extra peaks at 35.29° and 38.49° related to CuO phases were observed in $\text{Cu}_{6.62}\text{-SAPO-34}$ and $\text{Cu}_{8.84}\text{-SAPO-34}$ [30,31]. In the $\text{Cu}_x\text{-SAPO-34}$ with low Cu loading (<5.15 wt.%), no Cu-related phases were observed in the XRD patterns, suggesting that copper oxide was present in a highly dispersed amorphous state. The result further proved that too much CuO was present in the $\text{Cu}_x\text{-SAPO-34}$ catalysts with high Cu loadings. $\text{Cu}_{2.96}\text{-SAPO-34}$ sample exhibited the lowest crystalline nature mainly due to the mixture of SAPO-34 and amorphous material which was prepared at inappropriate pH value of 6.0.

3.3. The origin of the high hydrothermal stability of Cu-SAPO-34

3.3.1. $\text{NH}_3\text{-SCR}$ activity

The NO_x conversion over fresh and hydrothermally aged $\text{Cu}_{3.44}\text{-SAPO-34}$ is shown in Fig. 6(a). After 750 °C hydrothermal aging for 16 h, the SCR activity over the whole temperature range slightly increased, reaching ca. 90% at 200 °C, and maintained this level until 450 °C. Even after hydrothermal aging at 800 °C for 16 h, $\text{Cu}_{3.44}\text{-SAPO-34}$ still maintained excellent SCR activity, exceeding 90% from 225 to 400 °C. The SCR performance of fresh and hydrothermally aged $\text{Cu}_{5.15}\text{-SAPO-34}$ is shown in Fig. 6(b). The

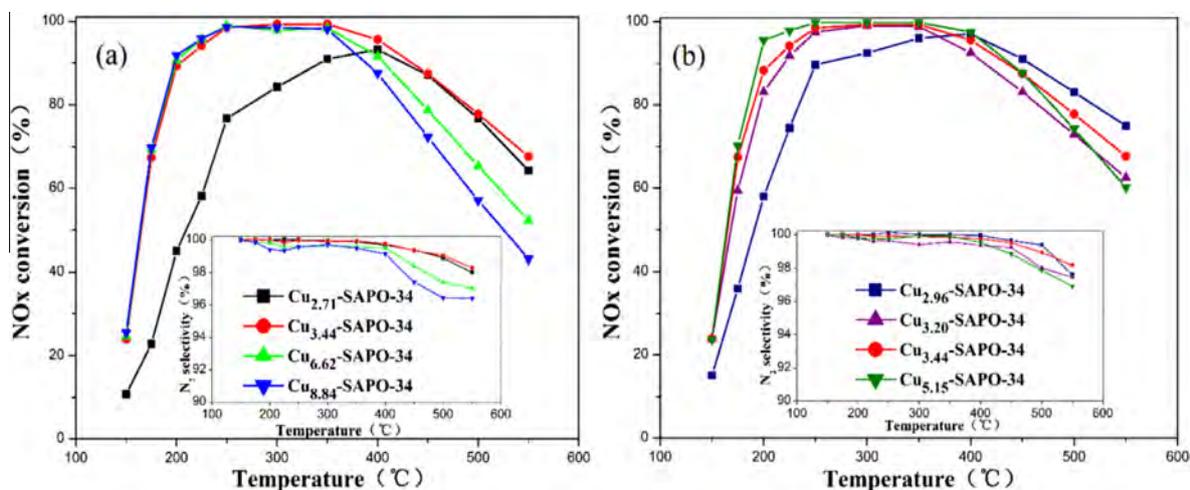


Fig. 4. NO_x conversion and N_2 selectivity over $\text{Cu}_x\text{-SAPO-34}$ samples.

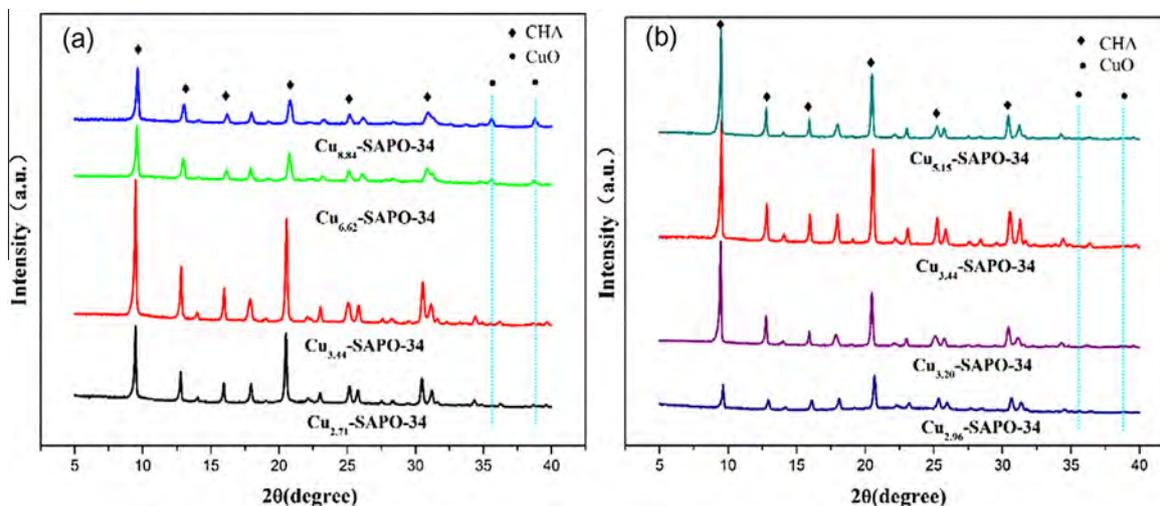


Fig. 5. XRD patterns of $\text{Cu}_x\text{-SAPO-34}$ samples.

results showed that the NO_x conversion above 350 °C decreased slightly over $\text{Cu}_{5.15}\text{-SAPO-34}$ after hydrothermal aging at 750 °C for 16 h, whereas hydrothermal aging at 800 °C for 16 h deactivated $\text{Cu}_{5.15}\text{-SAPO-34}$ completely. It was recognized that compared to the high Cu loading sample, the relatively low Cu loading sample was more resistant to relatively harsh hydrothermal treatment. It was speculated that high copper loading might cause the accumulation of copper species, forming CuO clusters, leading to structural instability.

3.3.2. $\text{H}_2\text{-TPR}$

The $\text{H}_2\text{-TPR}$ characterization was carried out to probe the distribution and reducibility of copper species. Fig. 7 shows that multiple Cu species coexist in the fresh and aged $\text{Cu}_{3.44}\text{-SAPO-34}$ and $\text{Cu}_{5.15}\text{-SAPO-34}$ samples. According to the literature [30,32–34], the low temperature and high temperature peaks were assigned to the $\text{Cu}^{2+} \rightarrow \text{Cu}^+$ and $\text{Cu}^+ \rightarrow \text{Cu}^0$ reduction steps, respectively. The reduction of CuO took place in a one-step mechanism ($\text{CuO} \rightarrow \text{Cu}^0$) and occurred in the same temperature range as the reduction of Cu^{2+} to Cu^+ (low temperature peak). Therefore, 210–250 °C was assigned to the reduction of isolated Cu^{2+} to Cu^+ , and 360–470 °C was attributed to the reduction of CuO to Cu^0 . The third peak at 670–770 °C was attributed to the reduction of Cu^+ to Cu^0 . EPR was an excellent technique for identifying the coordination environment of isolated Cu^{2+} ions, because all the other Cu species (CuO or Cu^+) were EPR silent [35,36]. By analyzing the hyperfine features of the hydrated $\text{Cu}_{3.44}\text{-SAPO-34}$ sample, $g// = 2.380$ and $A = 130.5$ G was obtained (Fig. S3). Zamadics et al. [35] reported that EPR signal ($g// = 2.381$, $A = 143$ G) in hydrated Cu-SAPO-34 could be assigned to Cu^{2+} located in the six-membered-ring as a complex $\text{Cu}(\text{O}_F)_3(\text{H}_2\text{O})_3$. The similar EPR signals for Cu-CHA catalysts reported by others [26,37] were also suggested to be this type of species, such as $g// = 2.39$ with $A = 111$ G

and $g// = 2.394$ with $A = 131$ G. Thus, this EPR signal should be due to the isolated Cu^{2+} located in the six-membered-ring for hydrated $\text{Cu}_{3.44}\text{-SAPO-34}$ catalyst. Through integrating the H_2 consumption peak areas, it was concluded that the first and the third reduction peaks were not equal for all samples, suggesting that the isolated Cu^{2+} ions might follow a one-step reduction mechanism ($\text{Cu}^{2+} \rightarrow \text{Cu}^+$). Some Cu^{2+} ions are too stable to be reduced to Cu^0 at temperatures below 900 °C. H_2 consumption peaks at 670–770 °C have been attributed to the reduction of Cu^+ to Cu^0 , either due to the Cu^+ formed from the first step of reduction of isolated Cu^{2+} or due to the Cu^+ originally existing in the framework of Cu-SAPO-34 [14,26].

What's more, the copper species distribution changed after hydrothermal aging over $\text{Cu}_{3.44}\text{-SAPO-34}$ and $\text{Cu}_{5.15}\text{-SAPO-34}$. It was reported that hydrothermal treatment of the Cu-SAPO-34 catalyst led to the migration of copper species [28,31,38], and the changes in copper oxidation states and its coordination environment directly affect the SCR activity. The $\text{H}_2\text{-TPR}$ results showed that ~6% CuO in $\text{Cu}_{3.44}\text{-SAPO-34}$ transformed to isolated Cu^{2+} after hydrothermal aging at 750 °C for 16 h (Table S1). As a result, the SCR activity increased over the aged $\text{Cu}_{3.44}\text{-SAPO-34}$ sample due to the presence of a larger amount of active Cu^{2+} ions. On the other hand, the copper species transformed from copper ions (Cu^{2+} and Cu^+) to CuO on the aged $\text{Cu}_{3.44}\text{-SAPO-34}$ (800 °C–16 h) and aged $\text{Cu}_{5.15}\text{-SAPO-34}$ (750 °C–16 h), leading to the decline of the SCR activity.

3.3.3. UV-vis-DRS

As shown in Fig. 8, UV-vis-DRS spectra were obtained to verify the nature of copper species of fresh and aged $\text{Cu}_{3.44}\text{-SAPO-34}$ samples. For fresh and aged $\text{Cu}_{3.44}\text{-SAPO-34}$ samples, the charge transfer band at around 230 nm can be assigned to the transfer from $\text{O}_{\text{SAPO-34}}$ to isolated $\text{Cu}^+/\text{Cu}^{2+}$ ions [39,40]. A broad band in

Table 4

The Cu species distribution on the $\text{Cu}_x\text{-SAPO-34}$ samples with different Cu loadings.

Catalysts	NO _x conversion at 200 °C, %	^a Cu ²⁺ , wt.%	NO _x conversion at 450 °C, %	^a CuO, wt.%
$\text{Cu}_{2.71}\text{-SAPO-34}$	45	1.35	87	0.35
$\text{Cu}_{3.44}\text{-SAPO-34}$	90	1.93	87	0.58
$\text{Cu}_{6.62}\text{-SAPO-34}$	91	3.55	78	1.45
$\text{Cu}_{8.84}\text{-SAPO-34}$	92	4.69	72	2.07

^a Calculated by the amount of H_2 consumption from $\text{H}_2\text{-TPR}$ results (Fig. S2).

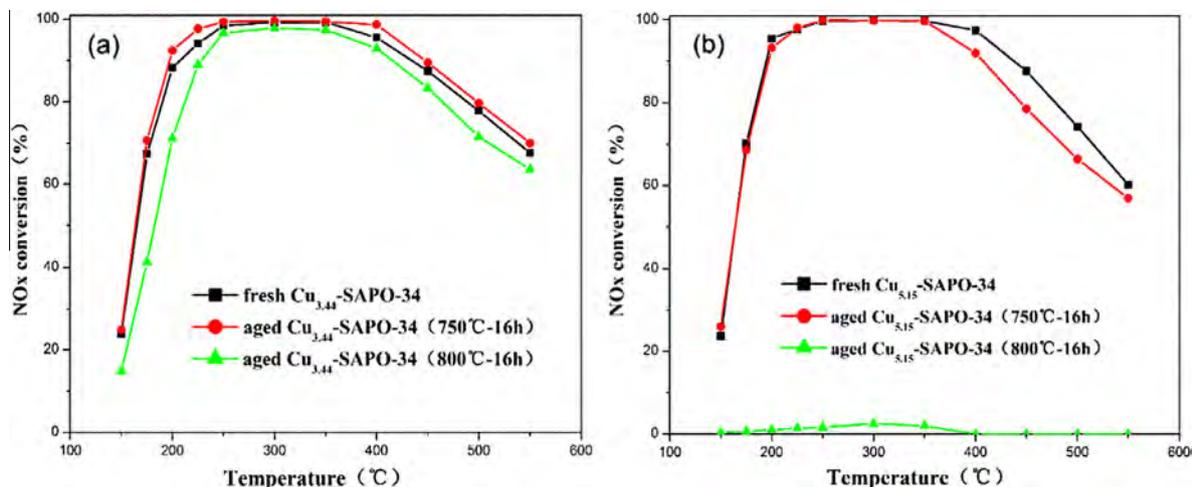


Fig. 6. NO_x conversion over the fresh and aged samples (a) Cu_{3.44}-SAPO-34 (b) Cu_{5.15}-SAPO-34.

the region of 600–800 nm was attributed to the electron d–d transitions of Cu²⁺ in distorted octahedral coordination surrounded by oxygen in CuO particles [31,41]. Compared with the results for the fresh sample, an increase in intensity of the band at 230 nm and a decrease in intensity of the band at 600–800 nm can be observed for 750 °C-aged Cu_{3.44}-SAPO-34. This indicated that the copper oxide species on the external surface of Cu_{3.44}-SAPO-34 could transform to copper ions and migrate into the cages of SAPO-34 during the hydrothermal process at 750 °C for 16 h, rather than aggregating to form larger clusters. In the case of the sample aged at 800 °C, the changes in the spectrum suggested the increase of CuO particles and decrease of isolated Cu²⁺ ions. The UV–vis–DRS results were in accordance with the H₂-TPR results.

3.3.4. XPS

The XPS results of the fresh and aged Cu_{3.44}-SAPO-34 samples are shown in Fig. 9. Cu 2p_{3/2} and Cu 2p_{1/2} peaks were observed in the ranges of 932.0–938.0 and 952.0–958.0 eV, respectively.

The Cu 2p_{3/2} peak around 935.8 eV as well as the satellite peak around 940.0–948.0 eV is used as a characteristic to determine Cu²⁺ [42,43]. The peaks of Cu 2p_{3/2} are further discussed in terms of the effect of hydrothermal treatments on the Cu valence state on the surface of the catalysts. It can be seen that the satellite peak around 940.0–948.0 eV was more obvious, indicating more isolated Cu²⁺ on the surface of Cu_{3.44}-SAPO-34 after hydrothermal aging at 750 °C which led to the performance improvement for NO_x removal to a certain degree. However, the peak at around 935.8 eV of aged Cu_{3.44}-SAPO-34 after aging at 800 °C decreased, indicating low Cu²⁺ concentration, which may be associated with the agglomeration of CuO [42,44] and in accordance with the H₂-TPR and UV–vis–DRS results, revealing the mechanism of deactivation during high temperature hydrothermal treatment.

3.3.5. Powder XRD

Fig. 10 shows the XRD patterns of the fresh and aged Cu_{3.44}-SAPO-34 and Cu_{5.15}-SAPO-34 samples. Except for the Cu_{5.15}-

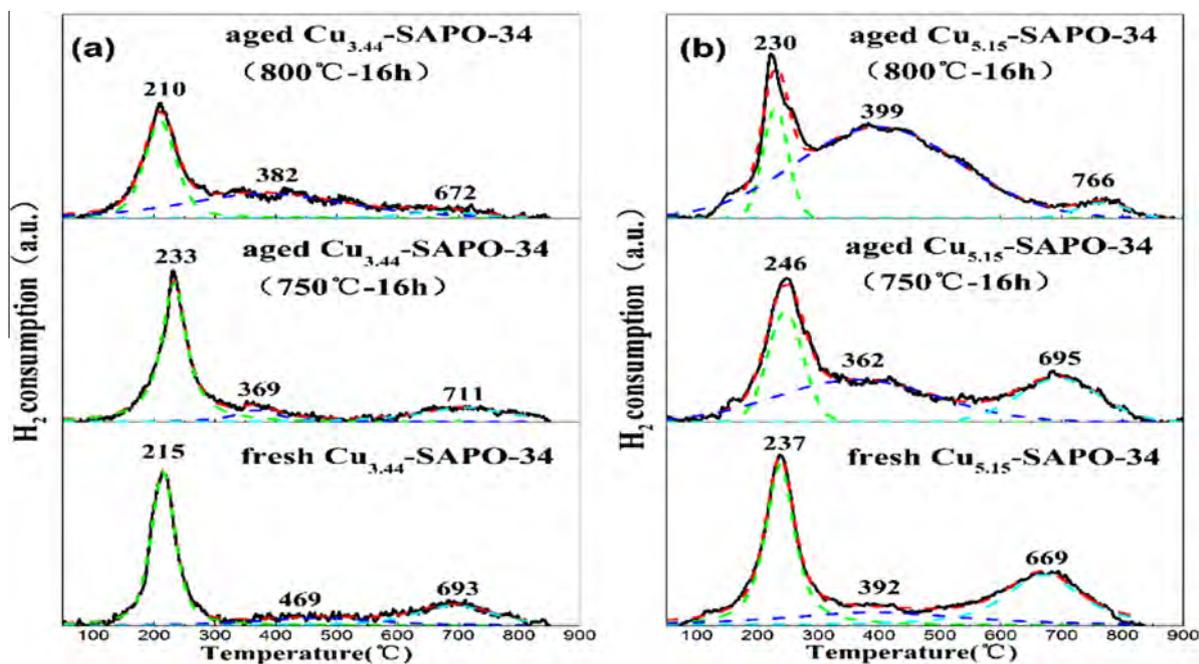


Fig. 7. H₂-TPR patterns of fresh and aged samples (a) Cu_{3.44}-SAPO-34 (b) Cu_{5.15}-SAPO-34.

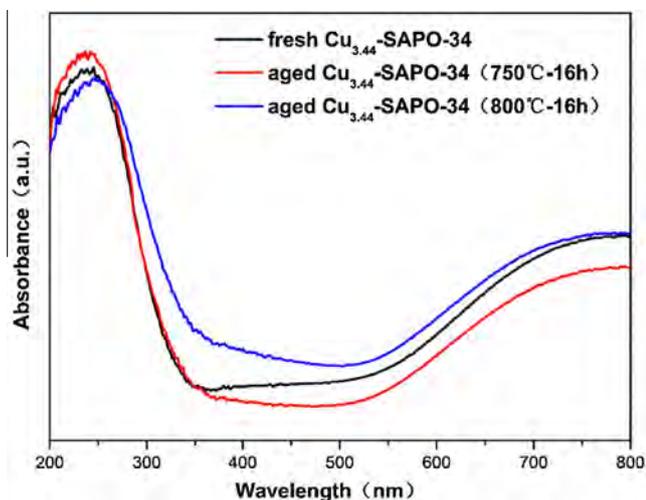


Fig. 8. UV-vis-DRS spectra of fresh and aged $\text{Cu}_{3.44}$ -SAPO-34 samples.

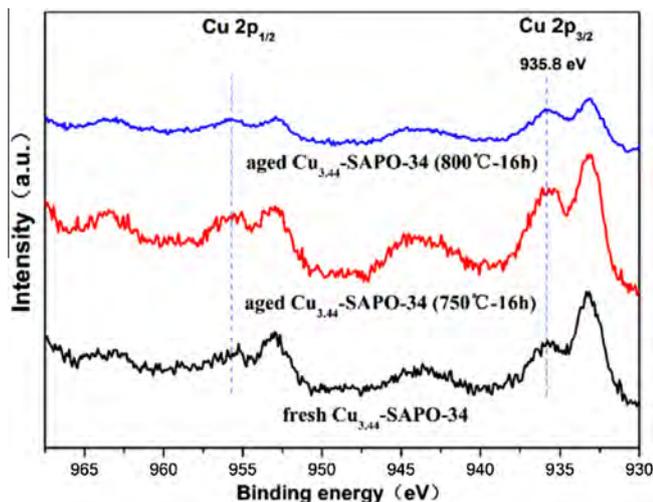


Fig. 9. XPS results of fresh and aged $\text{Cu}_{3.44}$ -SAPO-34 samples.

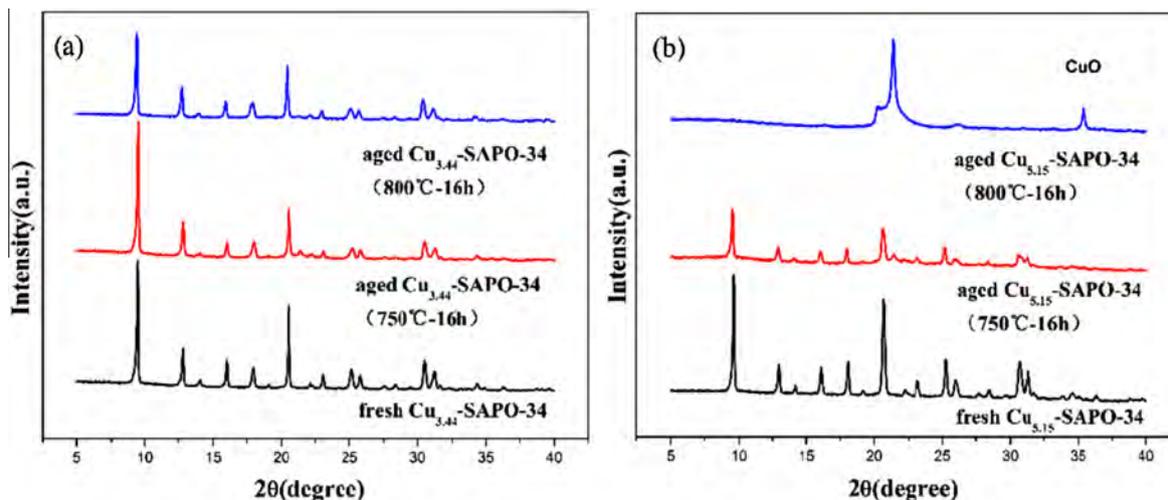


Fig. 10. XRD patterns of the fresh and aged samples (a) $\text{Cu}_{3.44}$ -SAPO-34 (b) $\text{Cu}_{5.15}$ -SAPO-34.

SAPO-34 sample aged at 800 °C for 16 h, all the other catalysts maintained the CHA zeolite structure. The collapse of the CHA framework resulted in the completely deactivation of the $\text{Cu}_{5.15}$ -SAPO-34 catalyst. On the contrary, hydrothermal treatment at 750 °C for 16 h had a negligible effect on the CHA structure of $\text{Cu}_{3.44}$ -SAPO-34, and the treatment under the higher temperature of 800 °C could not destroy the CHA structure. This was consistent with the moderate decrease in specific surface area and pore volume on aged $\text{Cu}_{3.44}$ -SAPO-34 samples measured by N_2 adsorption/desorption as shown in Table S2. On the other hand, the $\text{Cu}_{5.15}$ -SAPO-34 catalyst appeared to be more vulnerable by hydrothermal treatment. After the CHA structure collapsed, little pore structure was left on 800 °C-aged $\text{Cu}_{5.15}$ -SAPO-34. The Hydrothermal treatments indeed caused some damage to the physical structure probably due to the accumulation of copper species which had an adverse effect on the stability of the structure and led to lower NH_3 -SCR activity, but this effect was not very significant on the $\text{Cu}_{3.44}$ -SAPO-34 catalyst, indicating that this catalyst is a prospective candidate for industrial applications in the near future.

3.3.6. Acidity

To determine the density and strength of acid sites in the fresh and aged $\text{Cu}_{3.44}$ -SAPO-34 and $\text{Cu}_{5.15}$ -SAPO-34 catalysts, NH_3 -TPD measurements were carried out (Fig. 11). For the fresh $\text{Cu}_{3.44}$ -SAPO-34 sample, three NH_3 desorption peaks were observed. The peak (A) at 114 °C was found to be related to surface hydroxyl groups (Si-OH and P-OH). The peak (B) at 161 °C could be assigned to the weak structural Brønsted acid sites, whereas the peak (C) could be assigned to the strong structural Brønsted acid sites and Lewis acid sites (Si-OH-Al and Cu^{2+}) [14,45,46]. The distinction between acid sites present on the zeolite surface and those generated by deposition of copper can be realized by comparison of NH_3 -TPD results for Cu-SAPO-34 with different Cu loadings. As shown in Fig. S4, with the increase of Cu loading the absolute intensities of the peak between 230 and 320 °C increased and the absolute intensities of the peak between 320 and 500 °C decreased. Wang et al. [14] reported that Lewis acid sites can be created via Cu occupation of Brønsted acid sites in the pores. Wang et al. [47] also showed that the Cu^{2+} species can substitute for the proton (Si-OH-Al) of SAPO-34 supports. Hence, we deduced that the peak between 230 and 320 °C can represent the Lewis acid sites generated by deposition of copper, which increased with increasing Cu

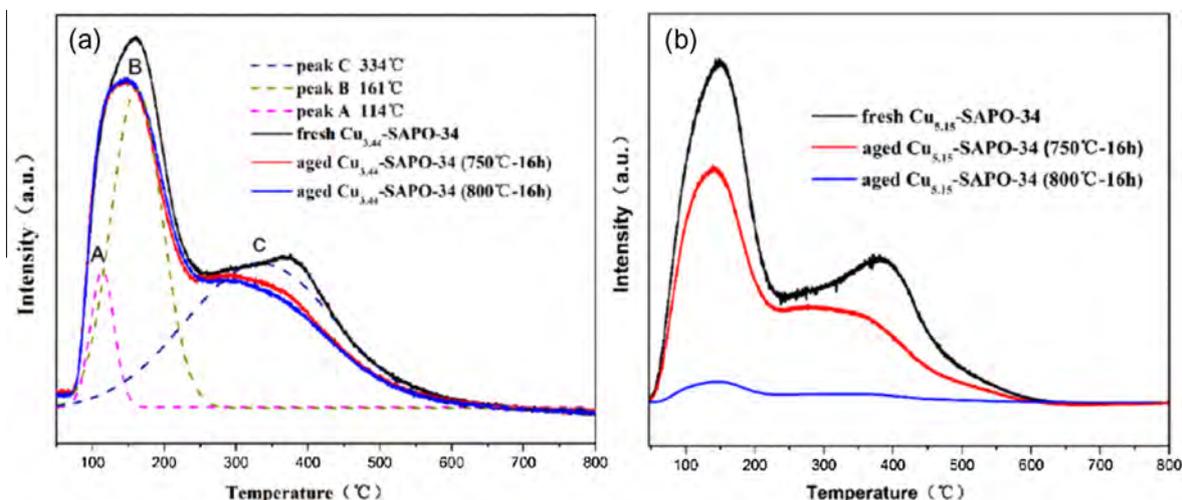


Fig. 11. NH_3 -TPD over fresh and aged $\text{Cu}_{3,44}$ -SAPO-34 and $\text{Cu}_{5,15}$ -SAPO-34 samples.

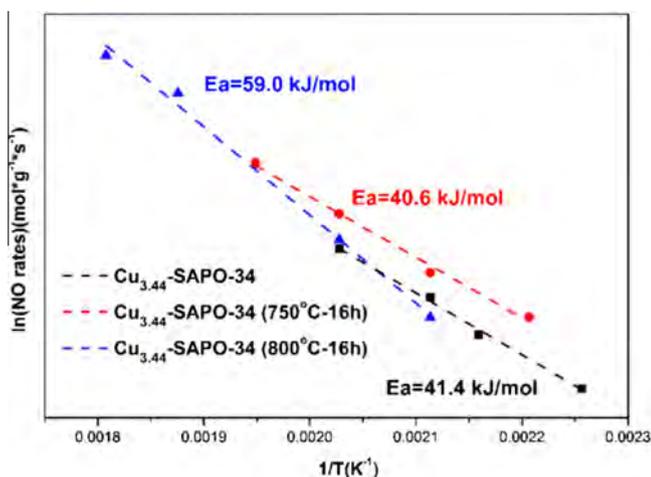


Fig. 12. Kinetic results of NH_3 -SCR over the fresh and aged $\text{Cu}_{3,44}$ -SAPO-34 samples.

loadings. The decline of the peak between 320 and 500 °C showed a reduction in the number of strong structural Brønsted acid sites. After harsh aging treatment, a slight decrease in the density of the acid sites was observed, indicating that the $\text{Cu}_{3,44}$ -SAPO-34 sample maintained most of the surface and structural hydroxyl groups. The Lewis acid sites generated by Cu^{2+} was little affected by hydrothermal treatment. However, aging treatment decreased the density of acid sites in $\text{Cu}_{5,15}$ -SAPO-34 more significantly than that in $\text{Cu}_{3,44}$ -SAPO-34. The acidity in $\text{Cu}_{5,15}$ -SAPO-34 after aging at 800 °C for 16 h were destroyed completely after such severe treatment.

3.3.7. Kinetic studies

In order to further prove the conclusions gained from the NH_3 -SCR tests, H_2 -TPR and NH_3 -TPD results, some kinetics studies were carried out over fresh and aged $\text{Cu}_{3,44}$ -SAPO-34 catalysts. The apparent activation energy (E_a) was estimated using the data obtained in the temperature range (170–280 °C) where NO_x conversions showed notable differences based on the Arrhenius plot of the rate (R) versus inverse temperature for NO (Fig. 12). The results revealed that hydrothermal treatment at 750 °C declined the E_a over $\text{Cu}_{3,44}$ -SAPO-34 from 41.4 to 40.6 kJ/mol. Otherwise, hydrothermal treatment at 800 °C increased the E_a over $\text{Cu}_{3,44}$ -

SAPO-34 from 41.4 to 59.0 kJ/mol. Therefore, the higher performance of NH_3 -SCR over the aged $\text{Cu}_{3,44}$ -SAPO-34 (750 °C–16 h) catalyst correlated well with its lower activation barrier. The hydrothermal treatment of 800 °C over $\text{Cu}_{3,44}$ -SAPO-34 increased the apparent activation energy, thereby decreasing the activity of NO_x reduction.

4. Conclusion

By using N_2 adsorption/desorption, XRD, H_2 -TPR, UV-vis-DRS, XPS and NH_3 -TPD, it can be concluded that the NH_3 -SCR activity and hydrothermal resistivity mainly depend on the framework pore properties, stability of the structure, copper species distribution and acidity of Cu-SAPO-34. With the increasing calcination temperature, the surface area, pore volume and structure crystallinity increased from 500 to 700 °C and then declined at 800 °C. When calcined at 700 °C, the Cu-SAPO-34 catalyst with copper loading ca. 3.44% performed optimal NH_3 -SCR activity. Additionally, >90% NO_x conversion can be maintained from 225 to 400 °C over $\text{Cu}_{3,44}$ -SAPO-34 catalyst after hydrothermal aging treatment at 800 °C for 16 h, indicating that $\text{Cu}_{3,44}$ -SAPO-34 catalyst is a promising candidate for industrial application.

Three types of copper species were observed on Cu-SAPO-34 sample, including isolated Cu^{2+} , CuO and Cu^+ ions. Hydrothermal aging at 750 °C led to the migration of CuO to isolated Cu^{2+} and an increase in the number of the active sites, which resulted in better NH_3 -SCR performance over the aged $\text{Cu}_{3,44}$ -SAPO-34 catalyst than the fresh catalyst. Meanwhile, the results of kinetics studies indicated that the apparent activation energy of aged $\text{Cu}_{3,44}$ -SAPO-34 (750 °C–16 h) was lower than the fresh sample. Therefore, the maintenance of isolated Cu^{2+} , the crystallinity and acidity as well as the low activation barrier were the origin of a high hydrothermal stability for the Cu-SAPO-34 catalyst.

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Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at <http://dx.doi.org/10.1016/j.cej.2016.02.086>.

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