

# CO oxidation on CuO/CeO<sub>2</sub> catalyst prepared by solvothermal synthesis: influence of catalyst activation temperature

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**Abstract** A series of CuO/CeO<sub>2</sub> catalysts were prepared using a solvothermal method and a subsequent activation process. The influences of activation temperature (300–600 °C) on physicochemical properties and catalytic behavior of the prepared CuO/CeO<sub>2</sub> catalysts have been investigated by XRD, SEM, Raman spectroscopy, *S*<sub>BET</sub> measurement, XPS surface analysis, H<sub>2</sub>-TPR, CO-TPD techniques, and CO oxidation testing. The catalyst activated at 450 °C was found to have the highest catalytic activity, which can be ascribed to its higher dispersion of Cu species, higher concentration of oxygen vacancies, and larger amount of more active lattice oxygen. The lower activity of the CuO/CeO<sub>2</sub> catalysts activated at lower (<450 °C) and higher (>450 °C) temperatures are attributed to the weaker interaction between CuO and CeO<sub>2</sub>, and to the sintering of CuO nanoparticles, respectively.

**Keywords** Solvothermal method · Activation temperature · CuO/CeO<sub>2</sub> catalysts · CO oxidation · Environment · Health effects

## Introduction

As a major air pollutant, CO is emitted from many industrial processes, transportation, and domestic activities. In order to reduce or even eliminate the toxic emission, catalytic oxidation of CO is considered to be the most promising method (Prasad and Rattan 2010; Chuang et al. 2012).

Currently, there are two main types of catalysts for catalytic oxidation of CO: one is the noble metal catalysts such as Pd, Pt, and Au; the other is base metal catalysts such as Cu, Co, and Mn. Although the noble metal catalysts exhibit a very high activity for CO oxidation, the high price and limited availability restrain their large-scale industrial applications (Prasad and Rattan 2010). Hence, more and more attention has been paid to base metal catalysts in order to find an alternative catalytic component to minimize using or even replace the noble metals. Among them, CuO/CeO<sub>2</sub> catalyst is considered to be the most effective due to the synergistic effect between CuO and CeO<sub>2</sub> (Ayastuy et al. 2009, 2010; Jia et al. 2010, 2012; Zagaynov 2015).

It has been well illustrated that the activity of CuO/CeO<sub>2</sub> catalyst for CO oxidation is greatly influenced by its preparation methods (Luo et al. 1997; Tang et al. 2004; Zhang et al. 2002). Currently, supported CuO/CeO<sub>2</sub> catalysts are usually prepared by impregnation (Jia et al. 2012; Luo et al. 1997; Tang et al. 2004) and deposition–precipitation (Tang et al. 2004, 2005) methods. Impregnation

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method possesses the merit of simplicity and convenience (Luo et al. 1997; Manzoli et al. 2005), but the CuO particles over the prepared CuO/CeO<sub>2</sub> catalysts tend to aggregate and lead to the poor dispersion of CuO, thus resulting in lower activity for CO oxidation (Tang et al. 2004). The CuO species of samples prepared by deposition–precipitation method are uniformly dispersed, but the preparation process is complex and it is easy for the impurities such as precipitants to get into the catalysts (Tang et al. 2004). Consequently, a variety of methods have been developed to obtain highly active CuO/CeO<sub>2</sub> catalyst including solvated metal atom impregnation (Zhang et al. 2002), laser vaporization and controlled condensation (Sundar and Deevi 2006), template method (Luo et al. 2007a; Qi et al. 2012; Zhu et al. 2008), modified sol–gel method (Luo et al. 2007b, c), combustion method (Avgouropoulos and Ioannides 2007), and flame spray pyrolysis (Kydd et al. 2009). However, these methods generally suffer from some harsh conditions, such as the use of bis (toluene) copper complex (Zhang et al. 2002), laser (Sundar and Deevi 2006), surfactants (Luo et al. 2007a; Qi et al. 2012), templates (Zhu et al. 2008), and hyperthermal treatment (Avgouropoulos and Ioannides 2007; Kydd et al. 2009; Luo et al. 2007b, c). So it is highly desirable to explore new simple and effective preparation methods (Qin et al. 2010). Compared with the above methods, the solvothermal process has advantages such as simple preparation and inexpensive operation conditions. Besides, the size and shape of the resulted nanoparticles are easily controlled by adjusting reaction temperature, time, and pH in this process (Lai et al. 2015).

Herein, a series of CuO/CeO<sub>2</sub> catalysts were prepared via a solvothermal method using ethylene glycol (EG) as a solvent and a subsequent activation process. The effect of activation temperature on the catalytic performance of the CuO/CeO<sub>2</sub> catalysts for CO oxidation was investigated. In order to clearly elucidate the influence of activation temperature, the physicochemical properties of the prepared catalysts were characterized in depth by means of XRD, SEM, Raman, BET surface area measurement, XPS surface analysis, H<sub>2</sub>-TPR, and CO-TPD techniques. Furthermore, the catalytic performance of the catalysts was correlated with the results of physicochemical characterization.

## Experimental

### Catalyst preparation

All chemicals were purchased from Shanghai Chemical Reagent Corporation (PR China) and used as received without further purification in this work. CuO/CeO<sub>2</sub> catalysts were prepared by a solvothermal method using EG as a solvent and a subsequent activation process. Commercial CeO<sub>2</sub> support was pretreated at 120 °C for 4 h before use to remove the impurity adsorbed on the surface. After that, the treated CeO<sub>2</sub> was added into an EG solution of Cu(NO<sub>3</sub>)<sub>2</sub>·3H<sub>2</sub>O and then reacted at 180 °C for 3 h in a Teflon-lined autoclave with continuous stirring. The suspension was allowed to cool down to room temperature (RT). The precipitate was separated by filtration, then repeatedly washed with absolute ethanol, and dried at 120 °C for ca. 12 h. Finally, the precursor was activated in a furnace under static air atmosphere at different temperatures (300–600 °C) for 4 h at a rate of 10 °C min<sup>-1</sup>. The obtained CuO/CeO<sub>2</sub> catalysts were named CC-300, CC-400, CC-450, CC-500, and CC-600. The CuO loading in all the catalysts expressed as the CuO/(CuO + CeO<sub>2</sub>) wt% ratio maintains as constant (10 wt%).

### Catalyst characterization

XRD measurements were conducted with a Rigaku Ultima IV using Cu K<sub>α</sub> radiation source generated at 30 kV and 25 mA. Two theta angles ranged from 10 to 90° with a speed of 4° per minute. The crystallite size of CuO was determined from full widths at half maxima of CuO (002) and CuO (111) peaks at 35.5° and 38.7°. The cell parameter values were calculated by standard cubic indexation method using the intensity of the CeO<sub>2</sub> (111) peak.

The surface morphology was observed by scanning electron microscopy (SEM, FEI, Quanta 200 FEG).

Raman spectra (at 4 cm<sup>-1</sup> resolution) was obtained with a DXR-Raman instrument (Thermo Fisher Scientific, American) using the 532 nm exciting line (200 mW beam), five scans for each spectrum.

N<sub>2</sub> adsorption–desorption isotherms were obtained at -196 °C on a Micrometrics ASAP-2020 adsorption apparatus, after all samples were degassed under vacuum at 200 °C for 10 h. The specific surface areas

were calculated from the linear part of the Brunauer–Emmett–Teller plot.

The chemical states of the elements were investigated by X-ray photoelectron spectroscopy (XPS, kratos Axis Ultra DLD spectrometer), using Al K $\alpha$  radiation. All the binding energy values were calibrated using C 1s = 284.6 eV as a reference.

Temperature-programmed reduction with H<sub>2</sub> (H<sub>2</sub>-TPR) was carried out using 10 vol% H<sub>2</sub>/N<sub>2</sub> as a reducing gas in a quartz microreactor. Approximately 0.02 g of a freshly calcined catalyst was placed on top of glass wool in the reactor. The outlet of the reactor was connected to a glass column packed with molecular sieve 5 Å in order to remove the moisture produced from reduction. The flow rate of the reducing gas was kept at 50 mL min<sup>-1</sup>, and the temperature was raised from 50 to 500 °C at a rate of 10 °C min<sup>-1</sup>. The consumption of H<sub>2</sub> was monitored by a thermal conductivity detector (TCD).

The CO adsorption property was measured by temperature-programmed desorption of CO. The catalyst (0.1 g) was pretreated under He flow (50 mL·min<sup>-1</sup>) with the temperature ramping from RT to 500 °C at a rate of 10 °C min<sup>-1</sup>, and then held at 500 °C for 40 min before being cooled down to RT in He flow. The next step was CO adsorption at RT for 30 min, and then the catalyst was swept again with He for 3 h. Lastly, the sample was heated in flowing He (50 mL min<sup>-1</sup>) up to 500 °C at a rate of 10 °C min<sup>-1</sup>, while the desorbed species was detected with a quadrupole mass spectrometer (QMS, Balzers OmniStar 200).

### Catalytic testing

The catalytic activity for CO oxidation was tested in a stainless steel tube microreactor (i.d. = 6 mm). Typically, 0.2 g of catalyst was placed between two layers of quartz wool. Gas mixtures containing 4 vol. % CO and 10 vol% O<sub>2</sub> in N<sub>2</sub> were passed through the catalyst bed at a flow rate of 30 mL min<sup>-1</sup>, corresponding to a space velocity of 9000 mL g<sup>-1</sup> h<sup>-1</sup>. The samples were heated in N<sub>2</sub> (50 mL min<sup>-1</sup>) at a rate of 10 °C min<sup>-1</sup> from RT to 200 °C and held at this temperature for 40 min in order to remove possible impurities. After cooling to RT in N<sub>2</sub>, the feed gas was introduced into the system. The temperature was increased stepwise (by 5–10 °C), and the catalyst was held for about 30 min at each fixed temperature to

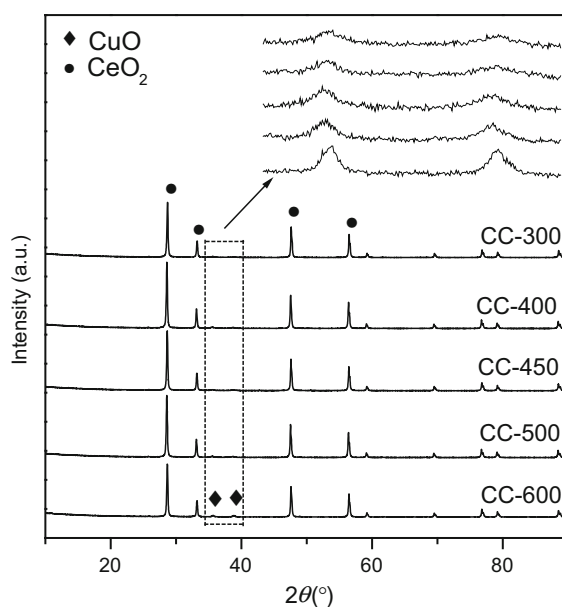
achieve steady-state activity. In this case, changes in the catalyst temperature due to heat release because of the possible occurrence of exothermic reactions of CO oxidation were not detected. To allow for the detection of CO and CO<sub>2</sub> with a flame ionization detector (FID), a methanator was inserted between one GC column and the FID. The gas compositions before and after the reaction were analyzed by an on-line gas chromatography with an FID. The activity was expressed by the conversion of CO.

## Results and discussion

### Characterization of catalysts

#### Structural and textural properties

XRD patterns of the CuO/CeO<sub>2</sub> catalysts are shown in Fig. 1. All samples exhibit strong diffraction peaks at  $2\theta = 29.2^\circ$ ,  $33.6^\circ$ ,  $48.6^\circ$ , and  $57.8^\circ$ , corresponding to cubic fluorite CeO<sub>2</sub> phase (JCPDS 34-0394). The narrow diffraction lines indicate better crystallinity of the CeO<sub>2</sub> support. Weak diffraction peaks corresponding to CuO appear at  $2\theta$  of  $35.5^\circ$  and  $38.7^\circ$  [JCPDS 41-0254] for all the catalysts, and the intensity of these peaks increases progressively with the increase in the activation temperature. The result indicates that the



**Fig. 1** XRD patterns of different CuO/CeO<sub>2</sub> catalysts

crystal size of CuO becomes larger as a result of particle sintering during the activation process at higher temperatures (Mai et al. 2011).

Table 1 shows the crystal sizes of CuO and CeO<sub>2</sub> on different CuO/CeO<sub>2</sub> catalysts estimated by the Scherrer equation. It can be seen that the crystal sizes of CuO enlarge with the increase in activation temperature from 300 to 600 °C, which can be connected to sample sintering. In addition, the crystal size of CeO<sub>2</sub> is essentially constant (39.0–40.6 nm) in all of the catalysts, which can be attributed to the commercial CeO<sub>2</sub> used in these experiments through the high-temperature roasting process.

The lattice parameters obtained from the CeO<sub>2</sub> (111) plane are also shown in Table 1. It is clear that lattice constriction has happened on all the CuO/CeO<sub>2</sub> catalysts comparing with support CeO<sub>2</sub>, indicating that some Cu<sup>2+</sup> ions have been incorporated into the CeO<sub>2</sub> lattice to form Cu–Ce–O solid solution in our catalysts (Cao et al. 2008; Guo et al. 2014). From Table 1, on the other hand, it is shown that the lattice parameter of CeO<sub>2</sub> varies in an irregular fashion as the activation temperature increases from 300 to 600 °C. This phenomenon can be attributed to the coexistence of two opposite effects (Fu et al. 2015): (1) the substitution of larger Ce<sup>4+</sup> (0.097 nm) by smaller Cu<sup>2+</sup> (0.072 nm) contracts the CeO<sub>2</sub> cell; (2) the formation of oxygen vacancies caused by the aliovalent doping (Ce<sup>4+</sup> by Cu<sup>2+</sup>) or creation of Ce<sup>3+</sup> (0.103 nm) sites expands the CeO<sub>2</sub> cell.

The SEM images of different CuO/CeO<sub>2</sub> samples calcined under different temperatures are shown in Fig. 2. From Fig. 2, we can see that the sizes of the CuO particles on the support surface increase with the increase in calcination temperature from 300 to 600 °C. In addition, the particles gradually

agglomerate when the calcination temperature rises from 300 to 600 °C.

Oxygen vacancies are considered as an indicator for the promotion of CO oxidation. Raman spectroscopy can provide direct evidence of oxygen vacancies and defects owing to changes in the vibrational structure of the CeO<sub>2</sub> lattice (Yen et al. 2012). Raman spectra of CeO<sub>2</sub> support and different CuO/CeO<sub>2</sub> catalysts are shown in Fig. 3. It can be seen that all the samples exhibit a strong vibration band at about 460 cm<sup>-1</sup>, corresponding to the F<sub>2g</sub> Raman vibration mode of fluorite CeO<sub>2</sub> (Kydd et al. 2009). Clearly, a red-shift of the F<sub>2g</sub> band is observed on all the CuO/CeO<sub>2</sub> catalysts compared with support CeO<sub>2</sub> (466 cm<sup>-1</sup>), and the reason may be the presence of oxygen vacancies, which is related to structural defects derived from partial incorporation of Cu<sup>2+</sup> into the CeO<sub>2</sub> lattice (Gnanakumar et al. 2014). This result is in good agreement with the decrease of lattice parameters obtained by XRD analysis.

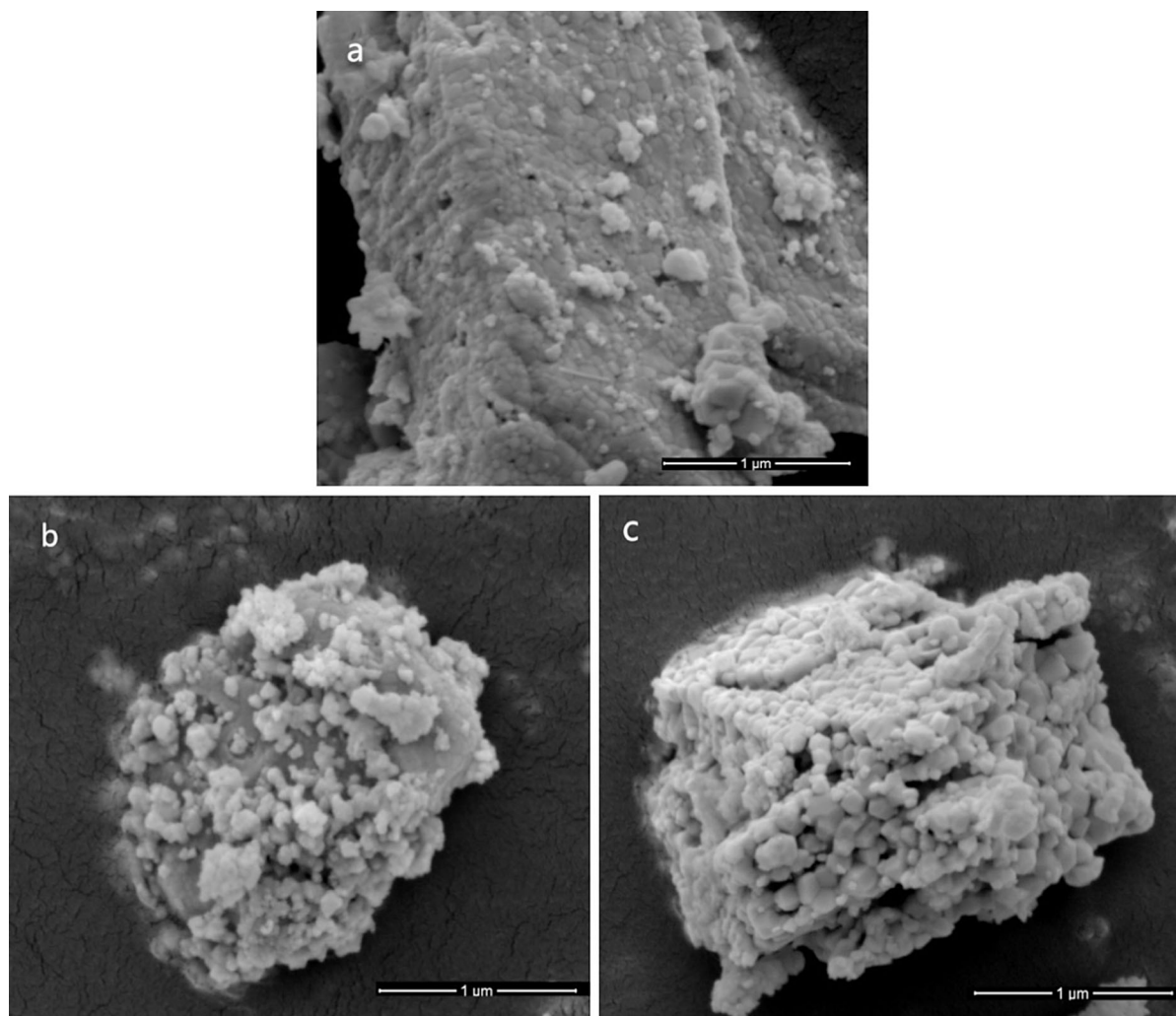
More specifically, the F<sub>2g</sub> bands shift from 462 to 455 cm<sup>-1</sup> when the activation temperature increases from 300 to 450 °C. However, the bands shift from 455 to 463 cm<sup>-1</sup> when the activation temperature further increases from 450 to 600 °C. Moreover, the width of the F<sub>2g</sub> band for the catalyst CC-450 is remarkably larger than those of the other catalysts. These results suggest that the catalyst CC-450 has the largest amount of oxygen vacancies among the investigated CuO/CeO<sub>2</sub> catalysts (Gnanakumar et al. 2014). The less amounts of oxygen vacancies over the catalysts activated at lower temperatures (<450 °C) can be attributed to the fact that lower temperature is unfavorable to the interaction between CuO and the support CeO<sub>2</sub> (Liu et al. 2013), while the less amounts of oxygen vacancies over the catalysts calcined at

**Table 1** Structural and textural properties of the CeO<sub>2</sub> support and CuO/CeO<sub>2</sub> catalysts

Sample	D <sub>CuO</sub> <sup>a</sup> (nm)	D <sub>CeO<sub>2</sub></sub> <sup>b</sup> (nm)	S <sub>BET</sub> (m <sup>2</sup> /g)	Lattice parameter <sup>b</sup> (nm)
CeO <sub>2</sub>	–	37.9	11.1	0.5414
CC-300	15.2	40.4	8.7	0.5386
CC-400	15.8	40.4	7.8	0.5405
CC-450	18.7	40.6	8.1	0.5397
CC-500	21.0	40.6	6.9	0.5408
CC-600	23.6	39.0	6.9	0.5393

<sup>a</sup> The particle size of CuO is the average of the calculated values based on CuO (002) and CuO (111)

<sup>b</sup> The particle size and lattice parameter of CeO<sub>2</sub> are based on CeO<sub>2</sub> (111) plane



**Fig. 2** SEM images of different samples: **a** CC-300, **b** CC-450, and **c** CC-600

higher temperatures ( $>450$  °C) can be attributed to the larger CuO particles formed at higher temperatures (Sun et al. 2015).

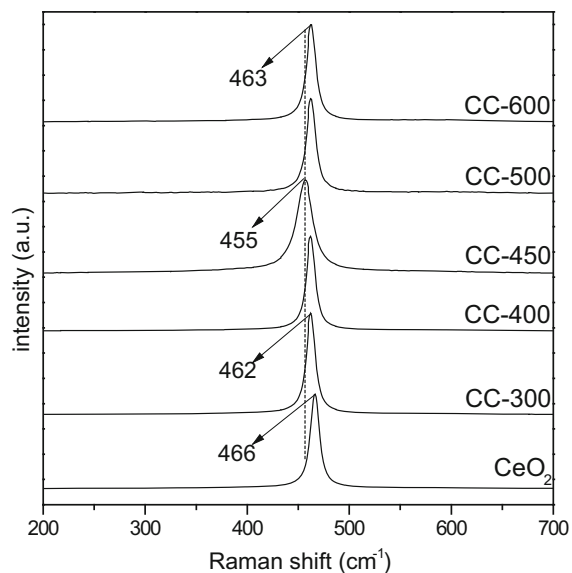
The  $S_{\text{BET}}$  values of the  $\text{CeO}_2$  support and CuO/ $\text{CeO}_2$  catalysts are listed in Table 1. All catalysts exhibit lower  $S_{\text{BET}}$  compared to  $\text{CeO}_2$  support, most probably due to the blocking of the pores of  $\text{CeO}_2$  by Cu species (Tang et al. 2004).

#### *Chemical state of elements*

In order to illuminate the chemical states of the elements existing in the CuO/ $\text{CeO}_2$  catalysts, XPS experiments were conducted. The Cu  $2p_{3/2}$ , Ce 3d, and

O 1s XPS spectra of the representative CuO/ $\text{CeO}_2$  catalysts are shown in Fig. 3a–c.

As shown in Fig. 4a, the broad peak of Cu  $2p_{3/2}$  binding energy at 930–937 eV can be deconvoluted into two peaks, centered at 934.4 and 932.7 eV, respectively. According to the literature (Av-gouropoulos and Ioannides 2003), the presence of higher Cu  $2p_{3/2}$  binding energy (about 934 eV) and the shake-up peak (about 938–945 eV) indicates the existence of  $\text{Cu}^{2+}$  species in these catalysts. Meanwhile, the lower Cu  $2p_{3/2}$  binding energy (932.7 eV) suggests the presence of reduced Cu species in these catalysts. On the other hand, the reduction degree of Cu species can be investigated by calculating the ratio



**Fig. 3** Raman patterns of the CeO<sub>2</sub> support and different CuO/CeO<sub>2</sub> catalysts

of the intensities of the satellite peaks to those of the principal peaks ( $I_{\text{sat}}/I_{\text{pp}}$ ), which is 0.55 for pure Cu<sup>2+</sup> (Avgouropoulos and Ioannides 2006). As shown in Table 2, the  $I_{\text{sat}}/I_{\text{pp}}$  values of these CuO/CeO<sub>2</sub> catalysts are lower than 0.55, confirming once again the presence of low-valence Cu species in these CuO/CeO<sub>2</sub> catalysts. The formation of the reduced Cu species may result from strong interaction of CuO with the support CeO<sub>2</sub> (Kundakovic and Flytzani-Stephanopoulos 1998), though the reduction of Cu<sup>2+</sup> under the procedure of XPS measurement cannot be completely discarded (Zhu et al. 2004).

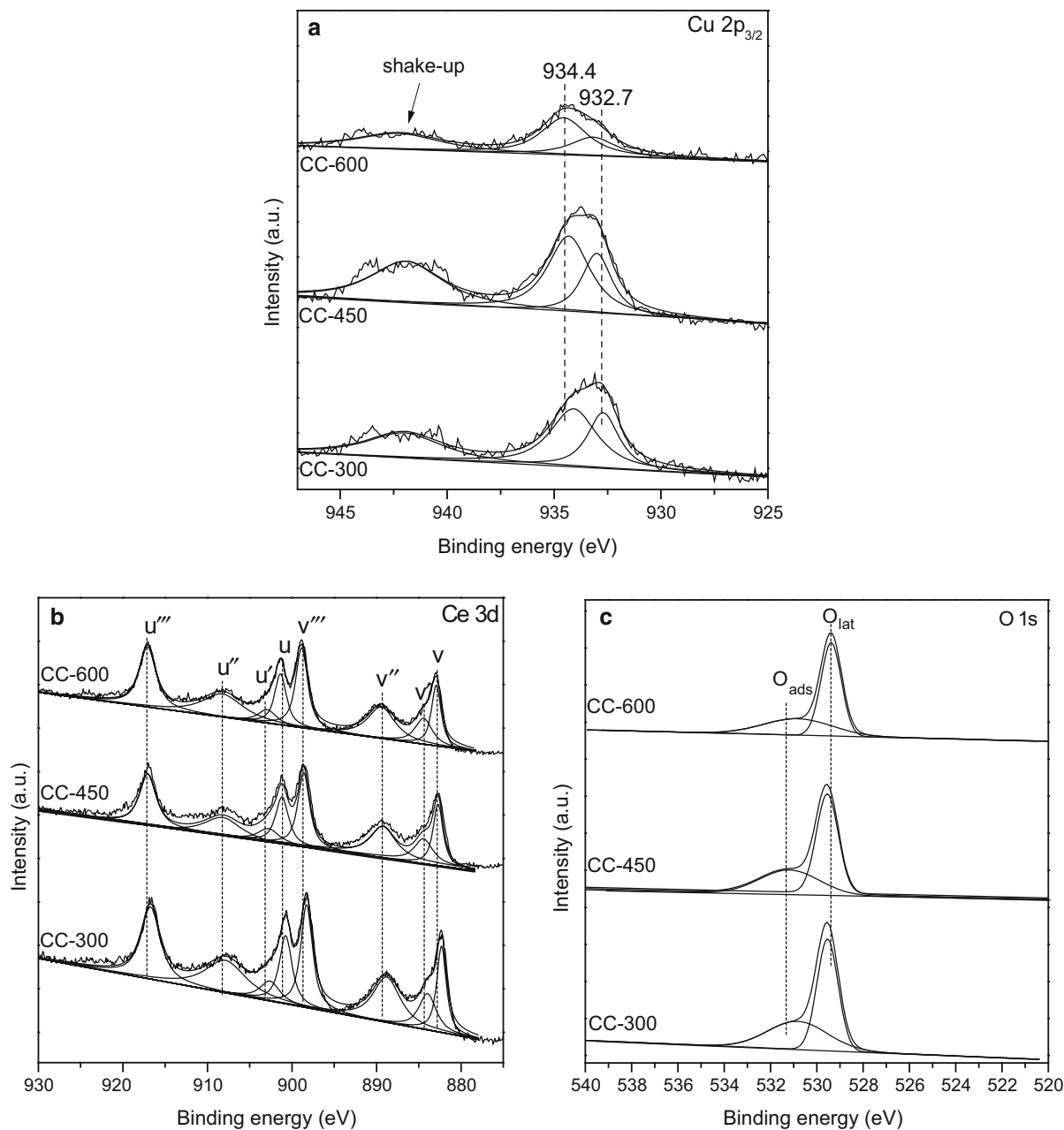
The Ce 3d XPS data obtained from different catalysts are shown in Fig. 4b. Two sets of spin-orbital multiplets relating to the 3d<sub>3/2</sub> and 3d<sub>5/2</sub> are labeled as u and v, respectively. According to previous research (Qi et al. 2012), the complex spectrum of Ce 3d is decomposed into ten components. Lines  $v_0$  and  $v'$  are contributed to the Ce 3d<sup>9</sup> 4f<sup>2</sup> O<sup>2</sup> p<sup>5</sup> and the Ce 3d<sup>9</sup> 4f<sup>1</sup> O<sup>2</sup> p<sup>6</sup> of Ce(III), respectively, and the same assignation can also be applied to the u structures. However,  $u_0$  and  $v_0$  are invisible in our samples, and the same phenomenon was also found over the CuO/CeO<sub>2</sub> catalyst prepared by the conventional impregnation method (Sun et al. 2015). Thus, an acceptable way to calculate the reduction degree of CeO<sub>2</sub> is

to consider the relative intensities of the  $u'$  and  $v'$  peaks to the total Ce 3d region.

$$\text{Ce(III)} (\%) = 100 \cdot \frac{[S(u') + S(v')]}{\sum [S(u) + (v)]}$$

The calculated percentages of Ce(III) for different CuO/CeO<sub>2</sub> catalysts are listed in Table 2. We can see that all the catalysts have the ability to carry out the reduction of CeO<sub>2</sub>, and the Ce(III) % value of the CC-450 catalyst is the largest. As reported, the existence of Ce<sup>3+</sup> in CeO<sub>2</sub> suggests the formation of oxygen vacancies (Li et al. 2011; Chang et al. 2006). Therefore, it can be inferred that the CC-450 catalyst has the highest level of oxygen vacancies, consistent with the above Raman results. Additionally, the values of Ce(III) % for these CuO/CeO<sub>2</sub> catalysts are approximately two times higher than that of pure CeO<sub>2</sub> reported in the literature (Ce(III) % = 6.75 %) (Cai et al. 2010), but the lattice parameters of these samples are similar to the reported values. This may be due to the fact that the sampling depth of XPS is lower than 10 nm, while the XRD analysis, from which the lattice parameters are determined, is a “bulk” technique. Therefore, there is a very large difference between the two kinds of analytical tools. In fact, almost all of the CeO<sub>2</sub> in bulk phase is still fully oxidized, thus the CeO<sub>2</sub> reduction level observed from XPS has a negligible effect on the lattice parameters. On the contrary, these CuO/CeO<sub>2</sub> catalysts show slightly lower lattice parameters caused by the incorporation of Cu<sup>2+</sup> into the CeO<sub>2</sub> lattice (Sun et al. 2015).

The XPS spectra of the O1 s region (Fig. 4c) show the contribution of two kinds of surface oxygen species. The low binding energy of 529–530 eV is characteristic of the lattice oxygen (denoted as O<sub>lat</sub>) associated with metal oxides, while the high binding energy in the region of 531–533 eV may be assigned to adsorbed oxygen (denoted as O<sub>ads</sub>) (Hu et al. 2009; Sun et al. 2014). The ratios of the peak area of lattice oxygen to total areas of lattice oxygen and adsorbed oxygen for these catalysts are presented in Table 2. It can be seen that the ratios for these CuO/CeO<sub>2</sub> catalysts increase with the increase in activation temperature, suggesting that a part of adsorbed oxygen has been transformed into lattice oxygen during the high-temperature activation process (Li et al. 2010).



**Fig. 4** XPS spectra of different CuO/CeO<sub>2</sub> catalysts: **a** Cu 2p<sub>3/2</sub>, **b** Ce 3d, and **c** O 1s

### Reducibility of catalysts

H<sub>2</sub>-TPR experiments were conducted to investigate the reducibility of the CuO/CeO<sub>2</sub> catalysts; the results are presented in Fig. 5. As shown, there are three peaks (denoted as  $\alpha$ ,  $\beta$ , and  $\gamma$ , respectively) being detected in these catalysts. A qualitative attribution of

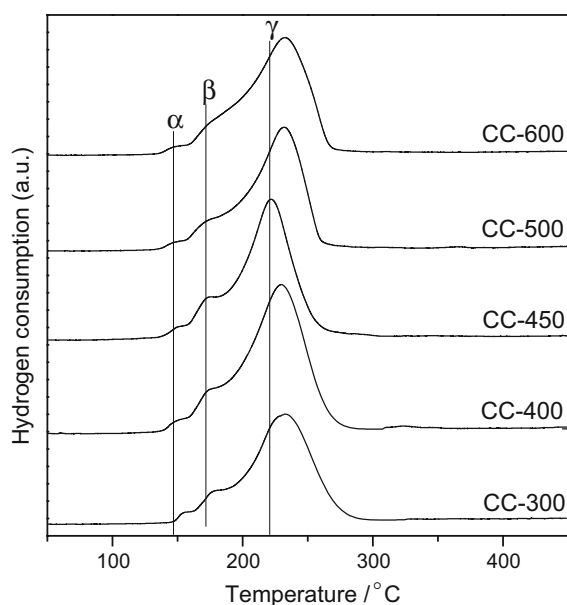
the TPR peaks to different Cu species over CeO<sub>2</sub> has been proposed by many researchers. For example, Luo and Avgouropoulos et al. (Avgouropoulos and Ioannides 2003; Luo et al. 1997) attributed the lower and higher temperature peaks to finely dispersed CuO strongly interacting with CeO<sub>2</sub> and larger CuO particles, respectively. Yen and Zhang et al. (Yen

**Table 2** XPS data measured for different CuO/CeO<sub>2</sub> catalysts

Catalyst	$I_{\text{sat}}/I_{\text{pp}}^{\text{a}}$	Ce (III) (%)	O <sub>lat</sub> <sup>b</sup> (%)
CC-300	0.505	12.9	58.4
CC-450	0.504	13.8	59.8
CC-600	0.504	12.2	62.9

<sup>a</sup> The ratio of the intensities of Cu 2p satellite peaks to those of the principal peaks

<sup>b</sup> The ratio of the peak area of lattice oxygen to total peak areas of lattice oxygen and absorbed oxygen

**Fig. 5** TPR profiles of different CuO/CeO<sub>2</sub> catalysts

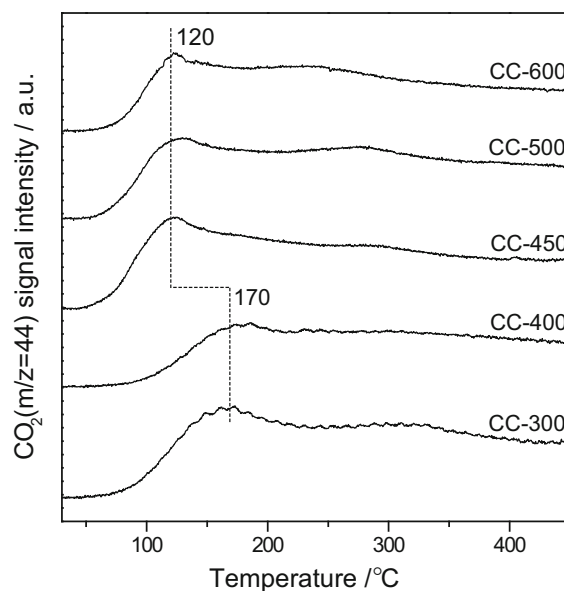
et al. 2012; Zhang et al. 2015) reported two reductive peaks which are attributed to the reduction of well-dispersed surface Cu species and Cu<sup>2+</sup> in the CeO<sub>2</sub> lattice, respectively. Zou and Moretti et al. (Moretti et al. 2011; Zou et al. 2006) suggested, for the more complex H<sub>2</sub>-TPR profiles, the contribution of both copper clusters and isolated Cu<sup>2+</sup> ions. In this work, based on these literature data and the above XRD results, the  $\alpha$ ,  $\beta$ , and  $\gamma$  peaks are attributed to the reduction of finely dispersed CuO strongly interacting with CeO<sub>2</sub>, Cu species doped into the support CeO<sub>2</sub>, and small two- and three-dimensional clusters of CuO weakly associated with the CeO<sub>2</sub> or the isolated bulk CuO without interaction with the CeO<sub>2</sub>, respectively.

As seen in Fig. 5, with the increase of activation temperature from 300 to 450 °C, the intensity of  $\alpha$  and

$\beta$  peaks increases slightly and then decreases noticeably with further increase of activation temperature from 450 to 600 °C. This result indicates that more Cu species is finely dispersed on the surface of CeO<sub>2</sub> or incorporated into the CeO<sub>2</sub> lattice for the CC-450 catalyst. On the other hand, the temperature of  $\alpha$  peak for the CC-450 catalyst is noticeably lower than that for the CC-300 catalyst. This result suggests that the interaction between the finely dispersed Cu species and CeO<sub>2</sub> for the former is stronger than that for the latter, since the stronger interaction leads to the lower reduction temperature (Scirè et al. 2010). Furthermore, the temperature of  $\gamma$  peak for the CC-450 catalyst is the lowest among these catalysts. This result suggests that the size of CuO clusters or bulk CuO particles on the CC-450 catalyst is the smallest, because the higher the reduction temperature is, the larger the CuO particle will be (Zhang et al. 2006). These results are in good agreement with those obtained by the above XRD and Raman analyses.

#### CO-TPD study

TPD profiles of CO<sub>2</sub> after CO adsorption on different CuO/CeO<sub>2</sub> catalysts are shown in Fig. 6. Almost all the adsorbed CO has been released after washing the sample under flowing He at RT. In the heating process, majority of adsorbed CO desorbs as CO<sub>2</sub> resulting

**Fig. 6** TPD profiles of CO<sub>2</sub> after CO adsorption at ambient temperature over different CuO/CeO<sub>2</sub> catalysts

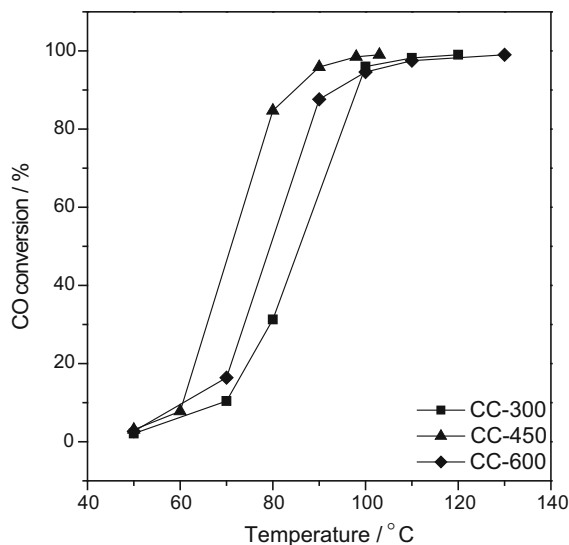


from different carbonate species produced by the reaction of adsorbed CO with lattice oxygen (Caputo et al. 2008). From Fig. 6, it can be seen that there are two CO<sub>2</sub> desorption peaks, including a main peak at 100–200 °C and a shoulder peak at >200 °C, which is consistent with the result of Avgouropoulos and Ioannides (Avgouropoulos and Ioannides 2008). According to the literature (Caputo et al. 2008), these two peaks may be related to different CO adsorption approaches: a fraction of CO has formed CO<sub>2</sub> at RT, which probably reacts with CeO<sub>2</sub> surface and adsorbs as carbonate, and releases at lower temperatures (100–200 °C); another fraction of the CO probably develops into bidentate carbonate on the reactive sites, which may evolve across intermediate species (may be as CO<sub>x</sub>) when the temperature increases and finally desorbs as CO<sub>2</sub> at higher temperatures (200–400 °C). More specifically, with the increase of activation temperature of the CuO/CeO<sub>2</sub> catalysts from 300 to 450 °C, the temperature of CO<sub>2</sub> desorption peak decreases noticeably from about 170 to 120 °C, implying that the produced carbonate species can desorb more easily (Luo et al. 1997). However, with further rise of the activation temperature, the CO<sub>2</sub> desorption peak temperature no longer decreases and the corresponding intensity of CO<sub>2</sub> desorption peak decreases slightly. As a result, it can be inferred that the CC-450 catalyst has the largest amount of more active lattice oxygen among the investigated CuO/CeO<sub>2</sub> catalysts.

#### Activity of catalysts

Figure 7 shows CO oxidation activities of CuO/CeO<sub>2</sub> catalysts (CC-400 and CC-500 have been removed for clarity). The T<sub>50</sub> (temperature for the CO conversion of 50 %) values are 86, 81, 71, 80, and 80 °C, for the catalysts CC-300, CC-400, CC-450, CC-500, and CC-600, respectively. It can be seen that the activity of CuO/CeO<sub>2</sub> catalysts for CO oxidation increases first with the increase of activation temperature, passing through a maximum at 450 °C, then decreases at higher activation temperatures.

There are many factors that can affect the activity of the CuO/CeO<sub>2</sub> catalyst for CO oxidation. On the basis of the above characterization results, the reason for the highest activity of the CC-450 catalyst can be explained as follows: (1) It is well known that the Cu species that are well dispersed and strongly



**Fig. 7** CO oxidation activities of different CuO/CeO<sub>2</sub> catalysts

interacting with the support CeO<sub>2</sub> are active toward CO oxidation, while larger bulk CuO particles contribute only a little or even negatively to the activity owing to the coverage of active species on the catalyst surface (Li et al. 2015; Luo et al. 1997; Mrabet et al. 2012). The above XRD result suggests that fewer larger bulk CuO particles are formed on the surface of CC-450 catalyst. In addition, the TPR result indicates that there are large amounts of well-dispersed Cu species strongly interacting with CeO<sub>2</sub> on the CC-450 catalyst than on the other catalysts. These structural features of the Cu species on the CC-450 catalyst are responsible for its higher activity. (2) CO oxidation on CuO/CeO<sub>2</sub> catalysts is considered to be a redox reaction involving oxygen vacancies. So, the higher the relative concentration of oxygen vacancies is, the better the performance of the catalyst will be (Jia et al. 2012; Li et al. 2010; Liu et al. 2010). Therefore, the highest activity of the CC-450 catalyst can be reasonably attributed to the highest concentration of oxygen vacancies obtained by Raman and XPS analyses. (3) It was reported that the oxidation of CO on metal oxides conforms to the following mechanism: (1) CO + O<sub>lat.</sub> → CO<sub>2</sub> + O<sub>vac.</sub>, (2) O<sub>2</sub> + 2O<sub>vac.</sub> → 2O<sub>lat.</sub>, where the O<sub>lat.</sub> and O<sub>vac.</sub> represent lattice oxygen and oxygen vacancy, respectively (Li et al. 2010). Obviously, the largest amount of more active lattice oxygen of the CC-450 catalyst as revealed by CO-TPD analysis accords well with its highest CO oxidation. Similar results were also

**Table 3** Comparison of CO oxidation activity of CuO/CeO<sub>2</sub> catalysts prepared by different methods

Catalyst	Preparation method	Reaction conditions	T <sub>100</sub> (°C)	Reference
Cu <sub>0.12</sub> Ce <sub>0.88</sub> O <sub>2</sub>	Surfactant template method	<i>sv</i> = 9600 mL/(gh), CO = 1 %	80 (T <sub>90</sub> )	Luo et al. (2007a)
Cu <sub>0.1</sub> Ce <sub>0.9</sub> O <sub>2</sub>	Modified sol–gel method	<i>sv</i> = 9600 mL/(gh), CO = 1 %	100 (T <sub>90</sub> )	Luo et al. (2007b)
Cu <sub>0.2</sub> Ce <sub>0.8</sub> O <sub>2</sub>	Hydrothermal method	<i>sv</i> = 16000 mL/(gh), CO = 1 %	120 (T <sub>90</sub> )	Mai et al. (2011)
Cu <sub>0.2</sub> Ce <sub>0.8</sub> O <sub>2</sub>	Co-precipitation+ hydrothermal treatment	<i>sv</i> = 30000 mL/(gh), CO = 1 %	180 (T <sub>90</sub> )	Li et al. (2015)
Cu <sub>0.19</sub> Ce <sub>0.81</sub> O <sub>2</sub>	KIT-6 template method	<i>sv</i> = 30000 mL/(gh), CO = 1 %	138 (T <sub>90</sub> )	Cui et al. (2014)
Cu <sub>0.19</sub> Ce <sub>0.81</sub> O <sub>2</sub>	Solvothermal method	<i>sv</i> = 9000 mL/(gh), CO = 4 %	105 (T <sub>90</sub> )	This work

reported by other researchers (Li et al. 2010; Liu et al. 2010; Zou et al. 2011).

The CO oxidation activities of the CuO/CeO<sub>2</sub> catalyst prepared by the present solvothermal method and those from the literature are presented in Table 3. It can be seen that our best-performing catalyst (CC-450) shows activities comparable to or even better than those of the CuO/CeO<sub>2</sub> catalysts prepared by other methods reported in the literature. It suggests that the use of simple solvothermal method can lead to appreciable activities, comparable to or even better than those achieved by other complex methods.

## Conclusions

A series of CuO/CeO<sub>2</sub> catalysts were prepared using a solvothermal method and subsequently activated at different temperatures. The catalyst activated at 450 °C shows the highest activity due to the better dispersion of CuO species as well as enhanced interaction between CuO and CeO<sub>2</sub>. The strong interaction between CuO and CeO<sub>2</sub> can induce the formation of large amounts of oxygen vacancies and activate the lattice oxygen, which is beneficial to the higher activity for CO oxidation. However, higher activation temperatures (>450 °C) lead to the aggregation and growth of CuO nanoparticles on CeO<sub>2</sub> surface due to the sintering, and the redundant larger bulk CuO species weaken the catalytic activity. On the other hand, the lower activity of CuO/CeO<sub>2</sub> catalysts activated at lower temperatures (<450 °C) can be attributed to the weaker interaction between CuO and the support CeO<sub>2</sub>.

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## Compliance with ethical standards

**Conflict of Interest** The authors declare that they have no conflict of interest.

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