Synthesis of cobalt doped mesoporous silica by using the rice husks as both silicon source and template with its catalytic oxidation of 2-methyl pyridine

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\textbf{Abstract:} Cobalt doped mesoporous silica (Co-SiO\textsubscript{2}) was prepared using the rice husks as both silicon source and template. Structural characterization of the catalysts was done by various techniques, such as X-ray diffraction, FT-IR, \textit{N\textsubscript{2}} adsorption/desorption, and scanning electron microscopy. Co-SiO\textsubscript{2} was used as a catalyst for the oxidation of 2-methyl pyridine and exhibited high substrate conversion (94.6\%) and good product (2-pyridinecarboxylic acid) selectivity (92.4\%). It even exhibited higher selectivity than Co-MTiO\textsubscript{2}, Co-MCM-41, Co-SBA-15. Fast hot catalyst filtration experiment proved that the catalyst acted as a heterogeneous one and it can be reused with almost the same activity.

\section{1 Introduction}

2-pyridinecarboxylic acid (2-PCA, picolinic acid) is an important intermediate for synthesis of physiologically active substances, dyes, and other synthetic chemicals\cite{1, 2}. In general, two basic methods have been employed for the synthesis of pyridinecarboxylic acids\cite{2}. One method is based on the hydrolysis of pyridinecarboxamides derived from pyridinecarbonitriles and the other is the oxidation of alkylpyridines/pyridineformaldehyde by air, nitric acid, selenium dioxide, etc.\cite{2-6}. However, these methods always need critical conditions and are still expensive, polluting and risky. Although several papers deal with the heterogeneous catalytic oxidation of 2-picoline\cite{2}, the selectivity for picolinic acid was only 19-22\% with 36-74\% conversion of 2-picoline. It is, therefore, of great practical interest to develop a more efficient, easily separable, reusable, and environment-friendly catalyst for the production of pyridinecarboxylic acid.

The mesoporous materials are of considerable interest in catalysis in recent years\cite{7-10}. The general synthetic strategies for mesoporous SiO\textsubscript{2} (such as MCM-41, SBA-15, etc.) have been developed by using surfactant as templates and tetra-ethyl orthosilicate as silicon source.

On the other hand, rice husk, a milling by-product of rice and also a major waste product of the agriculture industry. It is rich in silicon compound and has been used as silicon source to prepare silica long time ago\cite{11-14}. Recently, silica source from rice husk ash was used as an alternative for the synthesis of mesoporous silica\cite{15-16}. However, typical template, such as cetyltrimethylammonium bromide (CTAB) was still used as template in these synthesis processes. Whereas, the synthesis of porous silica using rice husk as templates was still very limited and in particular, the silica prepared by using rice husk as both silica source and templates has not been used in the oxidation of 2-methyl pyridine even though rice husk ash has been adopted as a catalyst support\cite{17}.

In the present work, rice husk were used as both silica source and template for the synthesis of cobalt doped mesoporous silica (Co-SiO\textsubscript{2}). It was used as a catalyst for the selectively catalytic oxidation of 2-methyl pyridine (2-MP) to 2-PCA acid for the first time.

\section{2 EXPERIMENTAL SEACTION}

\subsection{2.1 Synthesis of Co-SiO\textsubscript{2}}

Co-SiO\textsubscript{2} was synthesized with a process modified from Ref.\cite{15-16}: Rice husks (from a farm in Honghe, Yunnan province, China,) were rinsed in 1 mol/L HCl at 363 K for 24 h to eliminate metallic impurities, and filtered out and washed with distilled water until no Cl was detected. Then the husks were rinsed in 2-propanol repeatedly until the husks were completely dehydrated. The dehydrated husks were simply dipped in a closed vessel containing a 0.1mol/L solution of cobalt precursor (Co(NO\textsubscript{3})\textsubscript{2}-6H\textsubscript{2}O) in 2-propanol. The samples were shaken in an ultrasonic bath for 1 h to release the air bubbles emanating from the husks and allow the solution containing cobalt ion to enter the husks more easily, and then placed quietly for 24 h,
then the husks were filtered out. The treated rice husks were dried in air at room temperature. The final rice husks were calcined at 793.15 K for 12 h in an oven in air to burn off the organics. After naturally cooling to room temperature, a black product was obtained.

### 2.2 Characterizations

Pore size distributions, BET surface areas, and pore volumes were measured by nitrogen adsorption/desorption using a NOVAAtouch gas sorption analyzer (Quantachrome Corp.). Prior to the analysis, the samples were degassed at 150 °C for 1 h.

X-ray powder diffraction (XRD) experiments were conducted on a D/max-3B spectrometer with Cu Kα radiation, and scans were made in the 2θ range 0.1-5° with a scan rate 0.05°/min (low angle diffraction), and in the 2θ range 5-90° with a scan rate of 10°/min (wide angle diffraction).

Scanning electron microscopy (SEM) images were taken on a TM3000 Table Microscope at an accelerating voltage of 15 kV with the pressure in the sample chamber of 1 Torr.

FT-IR measurements were performed on a Thermo Nicolet 8700 instrument. Potassium bromide pellets containing 0.5% of the catalyst were used in FT-IR experiments and 32 scans were accumulated for each spectrum in transmission, at a spectral resolution of 4 cm⁻¹. The spectrum of dry KBr was taken for background subtraction.

### 2.3 Oxidation of 2-MP

The oxidation reaction was carried out at the atmospheric pressure as follows: A certain catalyst, 2-MP (AR, 0.5 g) and 10 mL solvent (acetic acid, acetic ether, water, acetone, acetonitrile, petroleum ether, chloroform, tetrahydrofuran, formic acid, dimethyl sulphoxide) were added successively into a 50 mL volumetric flask. Reaction mixture was heated to the set temperature. Reaction mixture was filtered under reduced pressure after the set time. Then residue was washed with CH₂OH and transferred into the 50 mL volumetric flask. Finally, it was analyzed by high performance liquid chromatography (Agilent 1200 series) with a packed column (Agilent ZORBAX SB C18 150x4.6 mm 5 μm) and an ultraviolet detector (diode array detector). The detector wavelength k = 254 nm and temperature of column was 313.15 K. The flow rate was 1 mL min⁻¹. The standard substances of 2-MP and 2-PCA were solvated in CH₂OH as standard solutions.

### 3 RESULTS AND DISCUSSION

#### 3.1 The characterization of catalyst

The wide-angle X-ray diffractogram patterns of Co-SiO₂, Co-MCM-41(CTAB) and the first recycled of Co-SiO₂ are shown in Figure 1.

As shown in Figure 1, it is worth noting that no distinct diffractions corresponding to any crystalline cobalt oxides are observed in the wide-angle diffractograms. This suggests the sample Co-SiO₂ is amorphous and also indicates that the cobalt oxides were well incorporated into the SiO₂ framework, or located only the walls of mesoporous materials. From Figure 1 it is also seen that the XRD patterns of the sample Co-SiO₂ and the sample first recycled of Co-SiO₂ show similar pattern to Co-MCM-41(CTAB), which prove them have similar framework to Co-MCM-41(CTAB)[7, 18]. This indicates that the framework of Co-SiO₂ has certain stability and does not change after the reaction. Moreover, there are observable lines corresponding to Co₃O₄ cubic phase in the XRD patterns of Co-SiO₂[18], but the lines disappear in the XRD patterns of the first recycled of Co-SiO₂. The phenomenon shows that Co₃O₄ can dissolve in the reaction, and indicates that Co₃O₄ is not incorporated into the SiO₂ framework, or located only the walls of mesoporous materials.

![Figure 1 XRD patterns of Co-SiO₂, Co-MCM-41(CTAB) and the first recycled of Co-SiO₂](image)

The N₂ adsorption/desorption isotherms and BJH pore-size distribution of Co-SiO₂ are shown in Figure 2.

![Figure 2 N₂ adsorption/desorption isotherms and corresponding pore size](image)
The isotherm of Co-SiO$_2$ can be classified as type IV according to the IUPAC convention and indicate the mesoporous structure of prepared materials[19,20]. This is also supported by the average pore width (5.3 nm), pore volume (0.29 cm$^3$/g), and BET surface areas (221m$^2$/g), which was much lower than that of Co-MCM-41. The decrease in surface areas could also be due to a coalescence process since sintering is favored by the presence of dopants[21]. The BJH pore size distribution of Co-SiO$_2$ (Inset of Figure2) also shows one primary pore size distribution in the mesopores region between 2.1 nm and 29nm and it indicates that the catalyst has irregular pore channels.

Fig. 3 show the SEM micrographs of Co-SiO$_2$(b), the template of rice husks(a), and Co-SiO$_2$ with different magnifications (c, d).

The non-uniform lamellar structure is different from that of typical MCM-41 prepared by TEOS as silicon source and surfactant templates[7]. And Co-SiO$_2$ is also different from MCM-48 prepared by using rice husk ash as silica source and CTAB as template. As shown in Figure 3a, the morphology of rice husk has two epidermis, and between inside and outside epidermis there is a interlayer[13, 14]. Figure 3b and Figure3c shows that the microstructure of Co-SiO$_2$ with pore system is well-distributed and compact, which elaborately duplicated the original hierarchical structure. The internal structure of Co-SiO$_2$ whice is composed of plates arranged in a crisscross pattern like loose honeycomb and contains a lot of square holes[14]. The topography of these holes shown in Figure 3d is in size of 8-20 µm.

The FT-IR spectra of Co-SiO$_2$ and Co-MCM-41 were recorded between 400 and 4000 cm$^{-1}$ and shown in Figure 4.

Both the samples exhibit the stretching vibration bands at around 460, 800 and 1100 cm$^{-1}$ for the tetrahedral SiO$_2$ structure[22]. Co-MCM-41 appears a very weak absorption peak at 968 cm$^{-1}$, and absorption peak at 960 cm$^{-1}$ disappears for Co-SiO$_2$, which implied that Si-OH groups were consumed and transformed to Si-O-Co bonds[23,24]. Furthermore, bands between the wavelengths of 2400 and 2300 cm$^{-1}$ that appear in both samples are assigned to the presence of CO$_2$ during the sample collection. In the hydroxyl region (3000-3500 cm$^{-1}$), the broad bands are observed at ca. 3400 cm$^{-1}$ and 1640 cm$^{-1}$ [25]. Especially, Co-SiO$_2$ appears two weak bands at 669 and 574 cm$^{-1}$, which is different from Co-MCM-41 and related to the vibrations of Co(III)-O bonds in Co$_3$O$_4$[26].

3.2 Catalytic performance

Using cobalt-doped SiO$_2$ as a catalyst, it was found that 2-PCA was the major product. For comparisons, we also studied the oxidation of 2-MP over undoped SiO$_2$, Co doped mesoporous silica (Co-MTiO$_2$[9], Co-MCM-41[7] and Co-SBA-15[23]). These comparisons are summarized in Table 1.

| Catalyst        | Conversion (wt %) | Selectivity (wt%) | TON |
|-----------------|------------------|------------------|-----|
| SiO$_2$         | <5               | <4               | --  |
| Co-SiO$_2$      | 94.6             | 92.4             | 313 |
| Co-MCM-41       | 77.6             | 83.9             | 256 |
| Co-SBA-15       | 81.5             | 82.6             | 269 |
| Co-MTiO$_2$     | 85.9             | 75.2             | 154 |
| First recycle   |                  |                  |     |
| (Co-SiO$_2$)    | 93.5             | 92.1             | 309 |
| Second recycle  |                  |                  |     |
| (Co-SiO$_2$)    | 92.5             | 89.3             | 305 |

Reaction condition: Substrate/oxidant, 0.33; reaction time, 6 h; reaction temperature, 363 K; catalyst, 50 mg; solvent, acetic acid.).
As expected, SiO₂ without doped by cobalt did not show any significant activity. It is apparent that the incorporation of cobalt into silica framework significantly enhanced the activity of reaction of 2-MP. Furthermore, even doped the same cobalt, Co-MTiO₂, Co-MCM-41 and Co-SBA-15 exhibited significantly lower selectivity and TON than Co-SiO₂. The conversion of 2-MP and selectivity of 2-PCA changed little by using Co-SiO₂ and the first recycled of Co-SiO₂ as catalyst, this may imply that the small amount of Co₃O₄ in Co-SiO₂ may have little influence on the oxidation of 2-MP. Although, Co-SBA-15 and Co-MCM-41 have higher surface areas (690.4 m²/g, 1041 m²/g, respectively), they are much less active than Co-SiO₂. This may imply that pore size would play more important role since the pore size of Co-SiO₂ (5.3 nm) is bigger than those of Co-SBA-15 (3.6 nm) and Co-MCM-41 (2.8 nm). In addition, porous structure diversity of Co-SiO₂ may be an important factor in reaction. This is because the structure-in-structure porous network could serve as transport paths for the distribution of electron onto the large surface of inner active mesoporous frameworks. Moreover, even compared with those catalysts used in the literatures [2-5], Co-SiO₂ with rice husk is more efficient since the yields of picolinic acid in these methods are very low (<20%).

3.3 Effect of solvents

The nature of solvents was known to have a major influence on reaction kinetics and product selectivity in the oxidation of 2-MP. Therefore, the effects of solvent on catalytic activity of Co-SiO₂ for the oxidation of 2-MP were studied first and summarized in Table 2.

| Solvent (10mL) | Conversion of 2-MP | Selectivity of 2-MP |
|---------------|-------------------|-------------------|
| Acetic acid   | 96.0              | 91.1              |
| Acetic ether  | 89.4              | 84.1              |
| Water         | 61.5              | 67.9              |
| Acetone       | 90.0              | 40.2              |
| Acetonitrile  | 87.9              | 83.8              |
| Petroleum ether | 95.7          | 0.9               |
| Chloroform    | 88.8              | 94.4              |
| TMF           | 64.5              | 67.3              |
| Formic acid   | 95.8              | 12.4              |
| DMSO          | 90.2              | 77.5              |

Reaction condition: Substrate/oxidant, 0.33; reaction time, 6 h; reaction temperature, 363 K; catalyst, 50 mg; solvent, acetic acid.

It was found that Co-SiO₂ catalyst had the best performance in acetic acid. In other solvents the catalyst exhibited significantly lower activities. This is due to possible partial decomposition of hydrogen peroxide because it was reported that the decomposition of H₂O₂ are faster in these solvent than acetic acid. The high efficiency of Co-SiO₂ because of the mesoporous materials are excellent activity for catalysis. It has been reported that the framework structure of Co-SiO₂ can impart dual catalytic activity to the catalyst and can form labile oxygen vacancies and the relatively high mobility of bulk oxygen species[8]. In addition, a complex with peroxy acetic acid was possibly formed in the pores of Co-SiO₂ which is relatively more hydrophobic and stable than hydrogen peroxide. Moreover, it has been reported that acetic acid does not only act as solvent, but also serves as a good oxidizing agent[27]. In short, the synergistic affects among doped cobalt, mesoporous framework of SiO₂, acetic acid, and hydrogen peroxide make Co-SiO₂ an effective catalyst for the oxidation of 2-MP.

3.4 Effect of reaction temperature

The dependence of the conversion of 2-MP on the reaction temperature was investigated in the temperature range 343-383K using over Co-SiO₂ is shown in Figure 5.

![Figure 5 Effect of reaction temperature on the conversion and selectivity over Co-SiO₂](image)

It is seen that the rate of 2-MP conversion decreases little with increase in reaction temperature while the selectivity of 2-PCA increases with increase in reaction temperature and passed through a maximum at 363 K. A further increase in the reaction temperature resulted in a slight decrease in the conversion but a fast decrease in the selectivity. This would be due to a quicker decomposition of H₂O₂ at higher temperature and the over oxidation that occurred with the rising temperature when 2-MP was almost completely converted after 363K. Considering the effect of temperature on both conversion of 2-MP and selectivity of 2-PCA, 363 K was chosen as the suitable temperature for the oxidation of 2-MP.

3.5 Effect of reaction time

The effect of reaction time on 2-MP reaction over Co-SiO₂ was investigated in Figure 6.
It is seen that the conversion of the 2-MP decreased little with time up to 12 h while the selectivity of 2-PCA increased with increase in reaction time and passed through a maximum at 6 h. Beyond this time, the selectivity of 2-PCA decreased with increase in reaction time. Considering the effect of time on both conversion of 2-MP and selectivity of 2-PCA, the optimum yield of 2-PCA could be achieved at about 6 h.

![Conversion and selectivity of 2-PCA over Co-SiO2](image)

**Figure 6** Effect of reaction time on the oxidation of 2-MP reaction over Co-SiO2. (Reaction condition: substrate/oxidant, 0.33; oxidant, H2O2; solvent, acetic acid; reaction temperature, 363 K; catalyst, 50 mg.)

### 3.6 Effect of catalyst concentration

Figure 7 presents the effect of catalyst concentration on conversion of 2-MP and selectivity of 2-PCA over Co-SiO2.

![Conversion and selectivity of 2-PCA over Co-SiO2](image)

**Figure 7** Effect of catalyst concentration on conversion of 2-MP and selectivity of 2-PCA over Co-SiO2. (Reaction condition: substrate/oxidant, 0.33; oxidant, H2O2; solvent, acetic acid; reaction temperature, 363 K; reaction time, 6 h.)

It is clear that the conversion of 2-MP had a slight fluctuations with the increase in the amount of catalyst, and this illustrates that the amount of catalyst changes has little effect on the conversion of 2-MP. Meanwhile the selectivity of 2-PCA slowly increased with increase in the amount of catalyst and passed through a maximum at 50 mg. Beyond this amount, the selectivity of 2-PCA decreased in the reaction mixture, and it is probable that large catalyst dosages provide excess active sites in the system to accelerate side reactions. Therefore, 50 mg would be the optimum catalyst amount used for the selectively producing 2-PCA. These results indicate that in the system catalyzed by Co-SiO2 only small amount of catalyst is active in the oxidation of 2-MP at higher amount of catalyst. This could be also attributed to the possible catalytic decomposition of H2O2 by Co-SiO2 and the competent interaction of metal oxo-species in Co-SiO2 sample with 2-MP, thus inhibiting the catalytic reaction.

### 3.7 The reusability and fast hot catalyst filtration experiment

In order to prove whether the catalyst is a heterogeneous one, experiments with fast hot catalyst filtration and studying the reactivity of the filtrate had been done by a modified process as literature[27]: 50 mg catalyst, 500 mg 2-MP, 20 mL HAc and 1 mL H2O2 was stirred at the temperature of 363 K for one hour, then quickly filtrate the catalyst. Let the filtration to react further with by adding 2 mL H2O2. The reaction carried out using this filtrate confirmed the absence of active metal ions as it exhibited conversion of 2-MP (16.9%) and the selectivity of 2-PCA (13.8%). This observation was also well supported by ICP-AES analysis of the filtrates obtained from the fast hot catalyst filtration where negligible amount of leaching of active cobalt species (0.005 wt.%) was detected. These indicated that the active component (cobalt) did not leach to the solution and the Co-SiO2 did act as a heterogeneous catalyst.

Finally, to check the stability and recycling ability as well as leaching of cobalt ions from Co-SiO2 under reaction conditions, recycling experiment were carried out using acetic acid as solvent. The typical recycling procedure was as follows: after the initial reaction, the catalyst was separated from mixture and washed with acetone and dried at 363 K, followed by the activation at 793 K for 2 h. The reaction was then carried out on the activated recycling catalyst. The results were also summarized in Table 1. The catalyst showed excellent reusability in the oxidation reactions. The major product was still 2-PCA, and the conversion of 2-MP and selectivity of 2-PCA changed little: the conversion from 93.5% at first use to 92.5% at second recycles, while the selectivity was from 92.1% to 89.3%, respectively. Obviously, the major product was still 2-PCA. After successive cycling of the catalyst, a minor change of catalytic activity was noticed. All the observations suggest that Co-SiO2 is a stable and highly effective catalyst for the oxidation of 2-MP.
4 Conclusions

In conclusion, cobalt-doped SiO₂ prepared by using the rice husks as both template and silicon source was an efficient catalyst for the oxidation of the 2-MP under relatively mild reaction conditions without adding any initiator. It even exhibited higher selectivity than Co-MTiO₂. Co-MCM-41 and Co-SBA-15. Fast hot catalyst filtration experiment proved that the catalyst acted as a heterogeneous one and it can be reused once with almost the same activity.

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