H$_3$PW$_{12}$O$_{40}$ Encapsulation by Nanoporous Metal Organic Framework HKUST-1: Synthesis, Characterization, Activity and Stability

Ezzat Rafiee$^{1,2,*}$ and Narges Nobakht$^2$

$^1$ Department of Inorganic Chemistry, Faculty of Chemistry, Razi University, Kermanshah, 67149, Iran
$^2$ Institute of Nano Science and NanoTechnology, Razi University, Kermanshah, 67149, Iran

* Corresponding author: E-mail: ezzat_rafiee@yahoo.com
Tel./fax: +98-833-427-4559

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Abstract

Hybrid composite material was obtained through encapsulation of H$_3$PW$_{12}$O$_{40}$ (PW) into HKUST-1 (Cu$_3$(BTC)$_2$, BTC = 1,3,5-benzenetricarboxylic acid), in molar composition of 5 Cu(NO$_3$)$_2$ · 3H$_2$O / 2.8 BTC / 0.3 PW / 0.6 CTAB by adding solutions of PW and copper salts to mixture of BTC and surfactant. The catalyst was characterized by various techniques including powder X-ray diffraction (XRD), Fourier transform infrared (FT-IR), scanning electron microscopy (SEM), energy dispersive X-ray (EDX), laser particle size analyzer, Brunauer Emmett-Teller (BET). The acidity of the catalyst was measured by a potentiometric titration with n-butylamine and PW/HKUST-1 presented very strong acidic sites with $E_i > 100$ mV. This nano catalyst was successfully used for the synthesis of various β-keto enol ethers at 45 °C with 51–98% yield after 5–75 min. The catalyst was easily recycled and reused at least four times without significant loss of its activity (94% yield after forth run). The presence of the PW in PW/HKUST-1 and reused PW/HKUST-1 structure, eliminating any doubt about collapse of the HKUST-1 after catalytic reaction and can be followed by FT-IR, XRD and SEM techniques. Brønsted and Lewis acidity of the PW/HKUST-1 catalyst was distinguished by studying the FT-IR and determined by chemisorption of pyridine. The strength and dispersion of the protons on PW/HKUST-1 was considerably high and active surface protons became more available for reactant.

Keywords: Heteropoly acid; Metal-organic framework; β-Keto enol ethers; Heterogeneous catalysts; Hybrid composite material.

1. Introduction

In the last decade, metal organic frameworks (MOFs) as one of the most important families of materials, constructed from metal containing nodes and organic linkers, has been considered as in several applications such as gas storage/separation and catalysis. The crystalline, porous structure in combination with huge surface area and pore volumes results in unique properties of these materials such as large porosity, high surface area, open metal centers, abundant aromatic ligands and ease of synthesis. Up to now, several reactions have been carried out using MOFs as solid Lewis acid catalysts or catalyst supports but MOF materials usually do not possess Brønsted acidity, therefore some attempts have been made to introduce such functionality into them. Recently, the use of conventional nano-metal functionality in combination with acidic supports such as Al$_2$O$_3$ and MOFs supports investigated in recent years. Heteropoly acids (HPAs) can act as excellent acid, redox, and bi-functional catalysts in a variety of synthetically useful selective transformations of organic substances, due to their strong Brønsted acidity and fast reversible multi-electron redox transformation activities. The excellent catalytic performance of HPAs qualifies them as prime candidates for the designed construction of multi-functional materials combining porosity and catalytic activity. However, applications of HPAs are limited by their low specific surface area and low stability under catalytic conditions. One of the strategies to overcome these drawbacks consists of their encapsulation within porous solid matrices such as MOFs. As a well-known MOF, a stable host HKUST-1,
can encapsulated various Keggin-type HPAs, and attractive catalytic performance can be endowed by these HPAs/HKUST-1.\textsuperscript{21–23} The opportunity of incorporating HPAs in porous MOFs arises as an attractive pathway to exploit the catalytic activity of these species and opens the opportunity to create new catalytic systems.\textsuperscript{24} Among the different protocols for the synthesis of useful building blocks, we selected the synthesis of \( \beta \)-keto enol ethers because of their very high impact as synths for the preparation of bioactive compounds, terpenoids, 4-alkylated-2-cyclohexenones, 2-aryl and 2-alkenyl-3-alkoxy-cyclohexenones and bicycle \( \alpha \)-octenones.\textsuperscript{25–27} However, the synthesis of \( \beta \)-keto enol ethers has received little attention despite their wide range of applications\textsuperscript{29–31} and developing a new, cost-effective and green protocol for this transformation is still a challenging task. The present work reports the use of \( \text{H}_3\text{PW}_{12}\text{O}_{40} \) on HKUST-1 (PW/HKUST-1) as an active and recyclable catalyst in the synthesis of \( \beta \)-keto enol ethers.

2. Experimental

2.1. Materials and Preliminary Characterization

Copper(II) nitrate trihydrate (99–104%), 1,3,5-benzene-tricarboxylic acid (BTC) (98%), cetyl trimethylammonium bromide (CTAB) (99+%), and PW (99+%) were obtained from Merck. Absolute ethanol (98%), dimedone (98%), n-butylamine (99%), ethyl acetate (98%), methanol (98%), tert-butanol (98%), cyclohexanol (99%), benzyl alcohol (99%), 4-methoxybenzyl alcohol (98%), 3-methoxybenzyl alcohol (98%), 2-nitrobenzyl alcohol (97%) and 2-propanol (99.5%) were obtained from Alfa-Aesar. 3-Methoxybenzyl alcohol (98%), 2-nitrobenzyl alcohol (99%), 4-methoxybenzyl alcohol (98%), n-butylamine (99%), absolute ethanol (98%), dimedone (98%), methanol (98%), n-butylamine (99%), ethyl acetate (98%), 1,3,5-benzene-tricarboxylic acid (BTC) (98%), cetyl trimethylammonium bromide (CTAB) (99+%), and PW (99+%) were obtained from Merck. Absolute ethanol (98%), dimedone (98%), n-butylamine (99%), ethyl acetate (98%), methanol (98%), tert-butanol (98%), cyclohexanol (99%), benzyl alcohol (99%), 4-methoxybenzyl alcohol (98%), 3-methoxybenzyl alcohol (98%), 2-nitrobenzyl alcohol (97%) and 2-propanol (99.5%) were obtained from Alfa-Aesar.

2.2. Preparation of the Catalyst

PW/HKUST-1 material was prepared from a synthesis mixture containing a molar composition of 5 Cu(NO\(_3\))\(_2\) · 3H\(_2\)O /2.8 BTC/ 0.3 PW/ 0.6 CTAB. For the synthesis of PW/HKUST-1, the mixture of BTC (0.21 g) and 0.07 g of CTAB in absolute ethanol (12 mL) was prepared, then PW (0.3 g) and Cu(NO\(_3\))\(_2\) · 3H\(_2\)O (0.4 g) were dissolved in distilled water (10 mL). Both solutions were combined and mixed under vigorous stirring (1000 rpm) for approximately 30 min at room temperature. The mixture was aged without stirring for 4 days at room temperature. The blue solid was then collected by filtration, washed with distilled water three times and dried at 60 °C for 24 h. CTAB was removed by Soxhlet extraction with ethanol performed for 24 h. The product was dried in air at 60 °C for 24 h.

2.3. Acidity Measurement

For the potentiometric titration, 0.05 g of solid was suspended in acetonitrile (90 mL) and stirred for 3 h. The suspension was titrated with a 0.05 mol/L solution of n-butylamine in acetonitrile. The potential variation was measured with a Hanna 302 pH meter using a double junction electrode.\textsuperscript{32}

2.4. Catalytic Reaction

A mixture of dimedone (1.0 mmol) and alcohol (4 mL) was stirred in the presence of the catalyst (0.06 g) at 45 °C. After completion of the reaction as indicated by TLC, the catalyst was separated by decantation (or filtration). The excess alcohol in the filtrate was removed by rotary evaporator and the crude purified by column chromatography over silica-gel (ethyl acetate/hexane, 1:4). All products were identified by comparing of their spectral data, \(^1\text{H}-\text{NMR} / ^{13}\text{C}-\text{NMR}, \) with those of the authentic samples.\textsuperscript{31,33–36}

3. Results and Discussion

3.1. Catalyst Characterization

The presence of the PW in the HKUST-1 framework was established by FT-IR (Fig. 1). FT-IR spectoscopies confirmed that the structures of HKUST-1 and the PW are retained in the composite PW/HKUST-1. The vibrational spectra of the composite exhibit the characteristic bands of both the HKUST-1 support and the Keggin structure of the phosphotungstate anion. In particular, the FT-IR spectrum of PW contains bands at 1080 cm\(^{-1}\) (PO\(_4\)), 982 cm\(^{-1}\) (W=O), and 892 and 796 cm\(^{-1}\) (W–O–W). For
PW/HKUST-1 the PO₄ and W=O vibration bands are shifted in comparison to bulk PW. This shift discloses the confinement effect of PW inside the porous solid.

As depicted in Fig. 2 (a, b), the SEM images of the PW/HKUST-1 samples show that the PW/HKUST-1 crystals have octahedral shape with the average size of approximately 300 nm. Fig. 2 (c, d) shows the N₂-adsorption–desorption isotherm and their respective BJH distribution of pore size graph. BJH pore size distribution of PW/HKUST-1 catalyst is presented in Fig. 2 (c) with average pore 3.8 nm. Fig. 2 (d) shows the sorption isotherms, can be classified as type IV (in the IUPAC classification), which is typical of mesoporous materials.

The low angle XRD patterns of the PW/HKUST-1 was also evaluated (Fig. 3). According to XRD pattern, synthesis method applied in this study yielded crystalline

Figure 1. FT-IR spectra of (a) PW, (b) PW/HKUST-1, (c) reused PW/HKUST-1, (d) PW/HKUST-1 after reaction with 0.1 mol of pyridine per gram of catalyst and (e) HKUST-1.

Figure 2. SEM images of PW/HKUST-1 (Scale bar: 1 μm), subfigure shows the more clear morphology of one particle (a), SEM images of reused PW/HKUST-1 (Scale bar: 1 μm) (b), BJH pore size distribution of PW/HKUST-1 (c) and N₂-adsorption–desorption isotherm (d).
PW/HKUST-1 materials and showed sharp reflections appearing at low angles (<5°/2θ) suggesting a high degree of mesoscopic ordering. XRD patterns together with the SEM images showed that the HKUST-1 was highly crystalline. The size distribution of the PW/HKUST-1 derived from a laser particle size analyzer, illustrates in Fig. 4. PW/HKUST-1 has a mean diameter of 190 nm and a wide size distribution with a polydispersity of 0.8. EDX is used to determine distribution of PW in the hybrid material by monitoring the contents of W and P. The EDX elemental mapping images of PW/HKUST-1 show highly dispersed PW is successfully incorporated in HKUST-1 and uniform distribution of P and W (from PW) and Cu (from HKUST-1) is also illustrate (Fig. 5).

![Figure 3. Low angle XRD patterns of PW/HKUST-1 (bottom) and reused PW/HKUST-1 (up).](image3)

![Figure 4. Grain size distribution of PW/HKUST-1.](image4)

The nature of the acid sites (Brönsted and Lewis) of the catalyst sample was determined by in situ FT-IR spectroscopy with chemisorbed pyridine. A calcinated powder sample in a sample holder was placed in specially designed cell. Then pyridine vapor was introduced under N₂ flow and the FT-IR spectrum was recorded.

Acidity of the PW/HKUST-1 catalyst was distinguished by studying the FT-IR spectrum of the catalyst after reaction with pyridine (Fig. 1(d)). Pyridine molecules were adsorbed on Lewis acid sites (1610 and 1450 cm⁻¹) and formed the pyridinium ion by interaction with Brönsted acid sites (1640 and 1540 cm⁻¹). Both types of adsorbed species contribute to the band at 1490 cm⁻¹. FT-IR spectrum of the catalyst shows contribution of pyridine adducts in the region 1400–1700 cm⁻¹. The formation of pyridinium ion was observed by absorptions at 1482, 1560 and 1605 cm⁻¹ (expanding region in Fig. 1). These results indicate that PW/HKUST-1 contains the strongest acid sites.

Also, the potentiometric titration curves obtained for HKUST-1, PW, PW/HKUST-1 are presented in Fig. 6.

![Figure 5. EDX elemental mapping images for the composite material PW/HKUST-1.](image5)
It is considered that the initial electrode potential \( (E_i) \) indicates the maximum strength of the acid sites and that the value from which the plateau is reached (mmol amine per g catalyst) indicates the total number of acid sites \( (n) \) that are present in the titrated solid. PW and PW/HKUST-1 presented very strong acidic sites according to the classification ranges \( (E_i > 100 \text{ mV} \) (very strong), \( 0 < E_i < 100 \text{ mV} \) (strong), \( -100 < E_i < 0 \text{ mV} \) (weak) and \( E_i < -100 \text{ mV} \) (very weak) and more acidic in comparison with HKUST-1. PW is well dispersed on the support surface and does not leach during the reaction. Therefore, a larger fraction of active sites are exposed to the surface, and this catalyst may exhibit excellent activity in organic reactions, even with a low catalyst loading. PW/HKUST-1 has an elemental composition comprising 39\% W, 16\% Cu, 17\% C, and 0.56\% P which was determined by ICP.

![Figure 6. Potentiometric titration curves of HKUST-1, PW, PW/HKUST-1.](image1)

3.2. Catalytic Reaction

The model reaction was carried out for studying the effect of catalyst loading and temperature on the product yield (Scheme 1). Initially, the quantity of the catalyst used in this reaction was optimized (Fig. 7). Influence of the amount of the catalyst on the model reaction was carried out at 45 °C when 4 ml methanol and 1 mmol dimedone were used and mixing rate was 500 rpm. Improvement in time of the reaction and yield was observed as the catalyst quantity increased from 0.02 to 0.06 g. Thus, 0.06 g of PW/HKUST-1 was the suitable choice for catalyst loading. Yield of the product was 64, 93, and 95\% when reaction temperature was 25, 45, and 60 °C respectively in the presence of 0.06 g of the catalyst and after 5 min. When the reaction temperature reached to 45 °C in the presence of 0.06 g of the catalyst the best result was obtained. According to the potentiometric titration results it seems that this yield was achieved per 0.084 mmol of active site of the catalyst \( (n = 1.4 \) total number of acidic sites; mmole acid site/g of the catalyst). When PW neat was used as catalyst in this optimized reaction conditions 94\% of the corresponding product was obtained after 5 min, but the catalyst is in homogenous phase and hard recovery of the catalyst is as a disadvantages of this method.

For comparison, the HKUST-1 material in the absence of PW, was used as a catalyst in the model reaction. The porous HKUST-1 itself shows 31\% yield after 20

![Figure 7. Effect of catalyst loading in the model reaction (4 ml methanol, 1 mmol dimedone, and mixing rate: 500 rpm at 45 °C).](image2)

Scheme 1. Model reaction for the synthesis of β-keto enol ethers using PW/HKUST-1
min. This observation revealed that Keggin PW species occluded in HKUST-1 pores significantly improved the catalytic activity of HKUST-1.

To evaluate the scope of the PW/HKUST-1 for the synthesis of β-keto enol ethers, different alcohols were used as reactants (Table 1). Primary, and secondary alcohols, reacted with dimedone without any significant difference to give the corresponding products in good to excellent yields. tert-Butanol was also reacted with dimedone to provide the corresponding β-keto enol ether in 50% yield (entry 6), while some previous approaches did not apply this alcohol or gave negative results for similar reactions.

Table 1 Synthesis of β-keto enol ethers using PW/HKUST-1 as catalyst.

| Entry | Alcohols | Time (min) | Yield (%) |
|-------|----------|------------|-----------|
| 1     | MeOH     | 5          | 98        |
| 2     | EtOH     | 10         | 98        |
| 3     | Propanol | 10         | 91        |
| 4     | Butanol  | 75         | 86        |
| 5     |          | 30         | 92        |
| 6     |          | 75         | 51        |
| 7     |          | 25         | 98        |
| 8     |          | 75         | 94        |
| 9     |          | 20         | 93        |
| 10    |          | 15         | 92        |
| 11    |          | 30         | 64        |
| 12    |          | 55         | 71        |

* Isolated yields, all products were identified by comparing their spectral data with those of the authentic samples.

3.3. Catalyst Recyclability

The recyclability of the PW/HKUST-1 was investigated to test the efficiency of this catalyst in consecutive cycles for the synthesis of β-keto enol ethers. The solid catalyst was recovered at the end of each reaction by simple filtration followed by washing with MeCN, dried at 60 °C and then reused in a fresh model reaction using the same experimental conditions. The results in Fig. 8 showed that synthesis of the product could reach 94% after being recycled for four times. It was observed that the catalytic system could be recycled at least four times with little decrease in catalytic activity and the amount of weight losing in this form of recovery was 5.5 wt.% but in continuous reaction system of course it should be considered more.

3.4. Stability of the Catalyst

The stability of the catalyst was evaluated by the characterization of the solid recovered after four consecutive cycles. Regarding the vibrational spectroscopy, the FT-IR spectra of the PW/HKUST-1 after catalytic reaction (Fig. 1(c)) are identical to the corresponding spectrum of fresh catalyst, displaying both the typical bands of HKUST-1 and the PW. The low angle XRD pattern of the reused PW/HKUST-1 exhibits an identical profile to the pattern of the composite before catalytic reaction (Fig. 3, up one). In fact, the main diffraction peaks of HKUST-1 are located at the same diffraction angles in both patterns, meaning that the crystalline structure of the HKUST-1 is retained after the catalytic cycles. Regarding the electronic microscopy studies, the morphology of the reused PW/HKUST-1 was similar to that of the corresponding composite of fresh catalyst and significant changes in the crystal morphology of the nanosized PW/HKUST-1 was not observed by SEM.

4. Conclusions

The PW/HKUST-1 composite material with average size of approximately 300 nm was prepared using simple
reaction technique. The acidity of the catalyst was measured by a potentiometric titration with n-butylamine and PW/HKUST-1 presented very strong acidic sites with $E_I$ > 100 mV. This nano catalyst was successfully used for the synthesis of various $\beta$-keto enol ethers at 45 °C with 51–98% yield after 5–75 min. The catalyst was easily recycled and reused at least four times without significant loss of its activity (94% yield after forth run). The presence of the PW in PW/HKUST-1 and reused PW/HKUST-1 structure, eliminating any doubt about collapse of the HKUST-1 after catalytic reaction was followed by FT-IR, XRD and SEM techniques. Brönsted and Lewis acidity of the PW/HKUST-1 catalyst was distinguished by studying the FT-IR. The strength and dispersion of the protons on PW/HKUST-1 was considerably high and active surface protons became more available for reactant.

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6. References

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Povzetek

Hibridni kompozitni material smo pripravili z enkapsulacijo $H_3PW_{12}O_{40}$ (PW) v HKUST-1 ($Cu_3(BTC)_2$, BTC=1,3,5-
benzenetrikarboksilan kislina), v molarjem razmerju $5 Cu(NO_3)_2$$/2.8 BTC/0.3 PW/0.6 CTAB$ z dodatkom raz-
topine $H_3PW_{12}O_{40}$ in soli bakra v znotraj BTC-ja in surfaktanta. Katalizator smo karakterizirali z naslednjimi tehni-
kami: rentgensko pravokovno difrakcijo (XRD), infrardečo spektroskopijo (FT-IR), vrstično elektronsko spektroskopijo
(SEM), energijsko disperzivno rentgensko spektroskopijo (EDX), laserskim analizatorjem za določanje velikosti delcev
in Brunauer Emmett-Teller metodo (BET). Kislost katalizatorja smo izmerili s potenciometrično titracijo z n-butilami-
nom in v materialu PW/HKUST-1 določili zelo kisla mesta z $E_i > 100$ mV. Tako pripravljen nanokatalizator smo us-
pešno uporabili za sintezo različnih $\beta$-keto enol etrov pri 45 °C z izkoristki od 51 % do 98 % in časom reakcij med 5 in
75 minut. Katalizator smo lahko reciklirali in ponovno uporabili brez znatnega zmanjšanja njegove aktivnosti (94 %
izkoristek po četrtem ciklu). Prisotnost PW v strukturi PW/HKUST-1 in v strukturi ponovno uporabljenega PW/HKUST-1 odpravlja vse dvome o kolapsu HKUST-1 po katalitični reakciji in jo lahko zasledujemo s FT-IR, XRD
in SEM tehnikami. Brønstedovo in Lewisovo kislost PW/HKUST-1 katalizatorja smo določili z FT-IR metodo in kemo-
sorbcijo piridina. Moč in disperzija protonov na PW/HKUST-1 je bila precej visoka, aktivna površina protonov pa je po-
stala dostopnejša za reaktante.