Quasi-HKUST Prepared via Postsynthetic Defect Engineering for Highly Improved Catalytic Conversion of 4-Nitrophenol

Minoo Bagheri, Arianna Melillo, Belen Ferrer, Mohammad Yaser Masoomi,* and Hermenegildo Garcia*

ACCESS | Metrics & More | Article Recommendations | Supporting Information

ABSTRACT: HKUST-1 [Cu$_3$(BTC)$_2$(H$_2$O)$_5$]$_n$nH$_2$OMeOH was submitted to thermolysis under controlled conditions at temperatures between 100 and 300 °C. This treatment resulted in partial ligand decarboxylation, generating coordinatively unsaturated Cu$^{2+}$ sites with extra porosity on the way to the transformation of the initial HKUST-1 framework to CuO. The obtained materials containing in part the HKUST-1 original crystal structure (quasi-MOFs) were used to promote 4-nitrophenol conversion to 4-aminophenol. Because of the partial linker decomposition, the quasi-MOF treated at 240 °C contains coordinatively unsaturated Cu$^{2+}$ ions distributed throughout the Q-HKUST lattice together with micro- and mesopores. These defects explain the excellent catalytic performance of QH-240 with an apparent rate constant of 1.02 × 10$^{-2}$ s$^{-1}$ in excess of NaBH$_4$ and an activity factor and half-life time of 51 s$^{-1}$g$^{-1}$ and 68 s, respectively, which is much better than that of the HKUST parent. Also, the induction period decreases from the order of minutes to seconds in the presence of the HKUST and QH-240 catalysts, respectively. Kinetic studies fit with the Langmuir–Hinshelwood theory in which both 4-nitrophenol and BH$_4^-$ should be adsorbed onto the catalyst surface. The values of the true rate constant ($k$), the adsorption constants of 4-nitrophenol and BH$_4^-$ ($K_{d,4NP}$ and $K_{d,BH4^-}$), as well as the activation energy are in agreement with a rate-determining step involving the reduction of 4-nitrophenol by the surface-bound hydrogen species.

KEYWORDS: heterogeneous catalysis, metal–organic frameworks, defect engineering, partial ligand removal, 4-nitrophenol reduction

INTRODUCTION

As it has been widely recognized, water pollution is a worldwide problem that greatly impacts living ecosystems. Nowadays, much research has been focused on the removal of water contaminants, such as aromatic pollutants. 4-Nitrophenol (4-NP) is a well-known toxic compound with high solubility and stability in water. It is a widely used synthetic intermediate in chemical processes to produce drugs, dyes, explosives, and pesticides. However, due to its biotoxicity, it is difficult to decompose by natural microbial degradation. Thus, the development of a cost-effective and ecofriendly method to remove this compound especially from agricultural and industrial wastewater, is of great importance.

The chemical reduction of 4-NP to 4-aminophenol (4-AP) can be a convenient strategy to remediate 4-NP wastes, since 4-AP is an important intermediate for the synthesis of analgesics and drugs, photographic developers, corrosion inhibitors, anticorrosion lubricants, and other specialty chemicals.

Several approaches have been applied for the reduction of 4-NP, including electrolytic reduction, metal-acid reduction, and catalytic hydrogenation. Electrolytic and metal-acid reduction reactions suffer from some disadvantages such as the need of an acidic or alkaline electrolyte, low efficiency, and poor selectivity. In comparison, the catalytic hydrogenation can be considered as a good prospective procedure to valorize 4-NP without generating any acidic waste, reaching high conversion and selectivity under mild conditions. Hence, recent studies have been more concerned with the efficiency of catalytic reduction of 4-NP into less toxic and valuable 4-AP, which is not only important from the environmental remediation point of view, but also beneficial as a valorization process. This catalytic reduction of 4-NP can be performed using sodium borohydride (NaBH$_4$), which is a mild reducing reagent, and the reaction can be conducted in aqueous medium under ambient conditions. Many studies have reported various catalytic systems. Among them, Cu-based complexes are convenient and efficient as Lewis acid catalysts combined with copper availability, economic affordability, and low toxicity. Therefore, several studies have reported on the catalytic
activity of copper compounds and copper oxides as catalysts for 4-NP reduction by NaBH₄.  

Metal–organic frameworks (MOFs), consisting of organic linkers and inorganic nodes defining a porous lattice, are among the most versatile heterogeneous catalysts. The active sites of MOFs can be coordinatively unsaturated metal ions, functional groups on the organic ligands, and/or guests inside the pores. Although there is a large number of MOFs with inherent catalytic properties, MOFs can also be used as supports, used as precursors of other materials, or combined with active components. Two methods are generally used to enhance the activity of MOFs as catalysts: (1) pore engineering to facilitate accessibility to the active sites and intracrystalline diffusion of substrates, reagent, and products, and (2) the generation of a coordinatively unsaturated position around metal sites by removal of solvent molecules. Pore engineering and structural defects have been generated during the synthesis by means of modulators that compete with the organic ligand during the formation of the coordinative bonds between the metal clusters and the linker. Alternatively, modification of porosity and generation of defects can be achieved by postsynthetic methods. Among them, the controlled thermal partial ligand decomposition has been proposed recently to create Lewis acid sites. The resulting materials are denoted as quasi-MOFs. It must be noted that defective MOFs have vacancies either of linkers or of metal nodes, whereas quasi-MOFs are materials with structures between the initial highly crystalline MOFs and the final metal oxides acquired by complete deligandation at relatively high temperatures. Control of the structural damage caused by heating of the MOF would allow one to prepare a new generation of materials with lower crystallinity, but with enhanced catalytic performance. The exact increase in activity that can be achieved following this strategy is still unknown, and it is of interest to determine which level of enhancement can be achieved before the material is finally transformed into the metal oxide.

The present study reports on the optimal quasi-Cu-MOF (HKUST-1) to catalyze the reduction of 4-NP by NaBH₄ as the reductant agent. Our strategy is based on thermal defect engineering to facilitate accessibility to the active sites and intracrystalline diffusion of substrates, reagent, and products. Pore engineering and structural defects have been generated during the synthesis by means of modulators that compete with the organic ligand during the formation of the coordinative bonds between the metal clusters and the linker. Alternatively, modification of porosity and generation of defects can be achieved by postsynthetic methods. Among them, the controlled thermal partial ligand decomposition has been proposed recently to create Lewis acid sites. The resulting materials are denoted as quasi-MOFs. It must be noted that defective MOFs have vacancies either of linkers or of metal nodes, whereas quasi-MOFs are materials with structures between the initial highly crystalline MOFs and the final metal oxides acquired by complete deligandation at relatively high temperatures. Control of the structural damage caused by heating of the MOF would allow one to prepare a new generation of materials with lower crystallinity, but with enhanced catalytic performance. The exact increase in activity that can be achieved following this strategy is still unknown, and it is of interest to determine which level of enhancement can be achieved before the material is finally transformed into the metal oxide.

The present study reports on the optimal quasi-Cu-MOF (HKUST-1) to catalyze the reduction of 4-NP by NaBH₄ as the reductant agent. Our strategy is based on thermal defect generation resulting in pore engineering and an extra high density of open metal sites in HKUST-1 with simultaneous distribution of both micro- and mesopores. These defects arise from the partial deligandation of the Q-HKUST network. Experimental evidence supports that controlled thermolysis can create highly active Lewis acid sites throughout the Q-HKUST. Contrary to the most widely reported use of metal oxides, HKUST-1 was performed in air at temperatures between 100 and 400 °C with heating at a 5 °C min⁻¹ rate for a period from 30 min to 2 h. The samples were denoted as QH-x, where “H” and “x” stand for HKUST and the deligandation temperature, respectively. Figure S1 shows the IR spectra of the resulting QH-x samples.

Catalytic Evaluation. The catalytic reduction of 4-NP was followed by UV–vis absorption spectroscopy. Before each run, the solutions were purged with N₂ to remove O₂. For this reason, 1.0 mg of catalyst was added to 25 mL of freshly prepared 4-NP (0.05 mM) and another 25 mL of NaBH₄ (10 mM) solutions at room temperature (rt). The solutions then were stirred vigorously. The conversion of 4-NP was subsequently determined by measuring the optical density at λₓₓₓₓ = 400 nm at periodic intervals. The catalytic 4-NP reduction performance was determined as follows:

\[ \% \text{reduction} = \frac{(C_i - C_f)}{C_i} \times 100 \]  

where \(C_i\) is the initial concentration of 4-NP and \(C_f\) is the concentration of 4-NP at any given time.

The kinetics of the reduction reaction can be obtained from eqs 2–4:

\[ \frac{-dC}{dt} = k_{app}C \]  

\[ \ln \left( \frac{C}{C_i} \right) = -k_{app}t \]  

\[ K = \frac{k_{app}}{m} \]

where \(k_{app}\) is the apparent reaction rate, \(m\) is the catalyst mass in gram, and \(K\) is the activity factor. To evaluate catalyst stability under the reaction conditions, the solid QH-x was isolated from the mixture by centrifugation and washed several times with deionized water and ethanol (1:1), and subsequently dried under vacuum at 100 °C for 10 h. The sample then was reused in the next reduction reaction. Four consecutive uses of the same sample were performed following the procedure mentioned above.

RESULTS AND DISCUSSION

HKUST-1 was obtained via solvothermal synthesis, by reacting copper(II) acetate and BTC ligand. The structure of HKUST-1 consists of 3D interconnected square-shaped pores (9 Å × 9 Å). The first report on the employ of quasi-MOFs in the catalytic reduction of 4-NP under moderate conditions.

The present study shows how the controlled partial thermolysis of HKUST-1 can produce the quasi-HKUST that exhibits a much enhanced catalytic activity for 4-NP reduction, as a consequence of the presence of unsaturated Lewis acid sites and the beneficial presence of micro- and mesopores. Q-HKUST has been prepared via partial deligandation in air at different temperatures in the range of 200–300 °C, with the highest catalytic performance for deligandation observed at 240 °C. This Q-HKUST is far more active than the parent pristine HKUST-1. The presence of residual phenyl rings of BTC in the mesopores facilitates 4-NP uptake via π–π stacking and hence increases the catalytic efficiency. Q-HKUST reusability and kinetics were also investigated.
Å) fabricated by connecting paddlewheel Cu(II) units through BTC ligands. Each Cu(II) metal ion with an octahedral configuration is coordinated to four different BTC linkers via oxygen atoms plus a Cu−Cu linkage (2.628 Å) and one H2O molecule coordinated in the direction of the Cu−Cu axis (Figure 1).

TGA analysis (Figure S2) revealed that thermal decomposition at moderate temperatures should cause partial deligandation of HKUST, generating additional pores and unsaturated inorganic metal centers. For this reason, HKUST-1 was heated at 120 °C for 14 h to expel guest molecules (H2O and EtOH) from the pores. Deligandation of the framework was performed by heating at 200, 240, and 260 °C for 30 min and 300 °C for 2 h or 30 min. Evolution of H2O and CO2 in the process was specified by MS analysis of the gases evolved in the head space upon heating. The corresponding samples are denoted, respectively, as QH-200, QH-240, QH-260, and QH-300.

The morphology of all samples was visualized by FE-SEM. The images revealed that parent HKUST-1 powder contains cubic nanoparticles. Thermal treatment decreases the average particle size, and particle agglomeration was observed (Figures S3–S12). EDS analysis shows the composition of the particles containing C, O, and Cu, while at temperatures of 400 °C only Cu and O can be detected, which indicated the complete conversion of HKUST-1 into CuO/Cu2O (Figure S12). In the case of QH-300, the presence of C is barely detectable by EDS (Figure S10).

PXRD of HKUST-1 and QH-x indicates that the original framework does not undergo significant changes up to 240 °C, except for a slight broadening of the peaks around 5−10° and diminution of their intensity, which indicate the initial stages of structural damage of HKUST. Upon increasing the treatment temperature to 260 °C, the characteristic peaks significantly decrease in intensity, eventually disappearing. For the 260 °C treatment, the presence of peaks corresponding to CuO/CuO (JCPDS nos. 05-0667 and 48-1548) was observed. When submitted to 300 °C, HKUST-1 is entirely transformed to CuO/Cu2O, implying the complete BTC decomposition (Figure 2).

The N2 adsorption data at 77 K for HKUST-1, QH-200, QH-240, and QH-260 display types I, (I, IV), IV, and V isotherms with BET specific surface areas of 1182, 913, 160, and 5.6 m2 g−1, respectively (Figure 3 and Table S1). Noticeable hysteresis in the adsorption of thermal decomposed samples implies the presence of large cavities. Moreover, the pore width distribution in QH-240 reveals a significant enhancement of the mesopores upon deligandation. The existence of micro- and mesopores in QH-240 can be ascribed to the formation of defects or vacant sites by partial deligandation with CO2 evolution and the development of a quasi-MOF structure. As a consequence of deligandation, a uniform distribution of coordinatively unsaturated Cu(II) ions in both micro- and mesopores should be present, and they can contribute to the improvement of the catalytic performance.

The XPS analysis was performed to gain insight into the distribution of Cu atoms among the various oxidation states after thermal treatment (Figure 4). In the XPS spectrum of HKUST-1, there are two peaks of Cu 2p3/2 and Cu 2p1/2 located at 933.6 and 953.5 eV, respectively, and the corresponding shakeup satellite of Cu 2p3/2 located at about 10 eV higher at 943.6 eV. All of these peaks agree with octahedral Cu2+.38,39 For QH-240, a new peak appeared at 932.8 eV attributed to the Cu+ species, accompanying a new intense peak at 935.2 eV that can be ascribed to the Cu2+ in tetrahedral coordination. These changes are associated with a considerable decrease in the peak intensities of octahedral Cu2+ (residual 8%).40 Thus, the XPS analysis confirms that a small fraction of Cu+ (8%) was created in QH-240 after
deligandation as well as the presence of Cu$^{2+}$ in tetrahedral sites (84%). In addition, the XP spectrum of QH-240 after reaction is almost similar to that of pristine QH-240. XPS data in combination with temperature-dependent PXRD showing that the final material after calcination at 300 °C is Cu$_2$O indicate the chemical reduction from Cu$^{2+}$ to Cu$^+$ in the thermal treatment. One possibility that cannot be ruled out at the moment is the reduction of Cu$^{2+}$ to Cu$^0$, followed by the subsequent Cu$^{2+}$ and Cu$^0$ comproportionation to Cu$^+$. The last possibility could be more likely for QH samples obtained at a higher temperature treatment, in which a massive formation of Cu$_2$O is observed.

Despite the dramatic changes shown in PXRD and surface area in the thermal treatment, vibrational spectroscopy that mostly reports on the functional groups of the organic ligand reflects minor variations, except for QH-300 and QH-400 for which the bands of the organic ligands have completely disappeared. Regarding the Raman spectra, the same bands, but much broader as determined by the width at half height, were recorded for HKUST-1 and QH-240. Figures S13 and

Figure 3. N$_2$ isotherm collected at 77 K and 1 bar and pore size distribution for HKUST, QH-200, QH-240, and QH-260.
S14 show the IR spectra of the samples under study and a comparison of the Raman spectra of HKUST-1 and QH-240. The EPR spectrum of HKUST-1 corresponds to Cu^{2+} in octahedral coordination. However, this characteristic EPR signal diminishes considerably in intensity to about a 10% in QH-240. Figure S15 shows the corresponding EPR spectra, thus reflecting the substantial changes occurring to the Cu^{2+} oxidation state and loss of the octahedral coordination in the thermal treatment.

Because of the large particle size, MOFs are not suitable for direct transmission electron microscopy. To observe the changes produced in the deligandation process, HKUST-1 and QH-240 were embedded in a polymeric matrix that allows one to cut thin slices by fast ion bombardment suitable for TEM imaging. The results are presented in Figure 5. As it can be seen there, while no particles are observed in the case of the HKUST-1 image that show a smooth internal surface, the QH-260 shows nanoparticles as well as the presence of macropores.

Catalytic 4-NP Reduction. The catalytic performance of the QH-x samples was examined for the reduction of 4-NP to 4-AP by screening some parameters on the catalyst treatment and reaction conditions, such as the deligandation temperature and catalyst mass. Kinetic data of reduction reaction such as activation energy, enthalpy, and entropy values of the adsorption on the most active catalyst were also determined.

By introducing NaBH₄ to the 4-NP solution, the strong absorption peak in the UV region shifted from 317 to 400 nm, which is associated with the creation of 4-nitrophenolate ions. Preliminary controls indicate that 4-NP reduction does not happen in the absence of catalyst even for a prolonged time. Upon addition of a proper amount of the suitable catalyst and as the reaction progressed, the absorption intensity at 400 nm gradually decreased, while a new peak appeared at 300 nm, indicating the production of 4-AP. Also, with the completion of the reduction reaction, the bright yellow solution became colorless (Figure S16).

**Influence of Deligandation Temperature.** The effect of temperature treatment on the catalytic activity of the material for 4-NP reduction was studied by comparing the performance of HKUST-1, QH-200, QH-240, QH-260, and QH-300 at room temperature (Table 1 and Figure 6). After QH-x catalysts were added, the peak at 400 nm sharply decreased and finally disappeared, while a new peak grows at 300 nm, demonstrating the conversion of 4-NP to 4-AP (Figure 6).

| Table 1. Influence of Deligandation Temperature on the NP Reduction Process Catalyzed by Various Quasi-MOFs |  
|---|---|---|---|---|
| quasi-HKUST | reduction time (min) | reduction (%) | induction period (s) |
| HKUST | 120 | 36 | 100 |
| QH-200 | 100 | 51 | 60 |
| QH-240 | 3 | 99.8 | 15 |
| QH-260 | 40 | 95 | 40 |
| QH-300 | 45 | 62 | 120 |
| QH-300-30 min | 45 | 65 | 125 |

*Experimental conditions: m(catalyst) = 1.0 mg, [4-NP] = 0.05 mM, [NaBH₄] = 10 mM, and T = 25 °C. QH-300-30 min was prepared as sample QH-300, but with 30 min of thermal treatment instead of 120 min.*
Agglomeration, and existence of both types of micro- and mesoporous structures are responsible for the enhancement of the catalytic activity. The relative catalytic activity of the HKUST-1 and QH-x samples also indicates that CuO (sample QH-300) is more active from the catalytic point of view than are framework octahedral Cu$^{2+}$ ions (HKUST-1), while the presence of Cu$_2$O (sample QH-260) increases the activity with respect to pure CuO. A QH-300 sample prepared with a shorter thermal treatment of 30 min instead of 120 min was also tested (Table 1) showing that a fast transformation of HKUST-1 into CuO takes place at this temperature. From Table 1, it can be concluded that tetrahedral Cu$^{2+}$ ions together with some minor Cu$^+$ ions prevalent in QH-240 are the most active sites for 4-NP reduction.

As it can be seen in Table 1, the catalytic performance is enhanced by increasing the deligandation temperature up to 240 °C and decreasing afterward. QH-200 and QH-260 catalysts were less active than the QH-240 catalyst because the concentration of Lewis acid sites in QH-240 was found to be higher than that of the other catalysts, while at the same time QH-240 exhibits a distribution of both micro- and mesopore structures (Figure 3).

The deligandation of HKUST up to 200 °C mostly leads to the removal of solvent molecules without any salient change in the framework porous structure, thus resulting in a moderate increase in catalytic activity. The concentration of open metal sites increases in parallel with the mesopore volume of the framework as deligandation continues up to 240 °C, which causes a notable enhancement in the catalytic 4-NP reduction, requiring a much shorter time to achieve almost complete conversion. Upon further temperature increase, most of the framework collapses, resulting in a dramatic decline in surface area and, therefore, in catalytic activity.

Also, the induction period ($t_0$), in which no catalytic reduction happens, decreases from 100 to 15 s in the presence of HKUST and QH-240 catalysts, respectively, while increasing afterward. Here, the induction time is attributed to the generation of the active reducing agent on the catalyst sites, and the 4-NP reduction reaction starts after this time. Therefore, the QH-240 sample was chosen as the best catalyst for additional studies on the catalytic 4-NP reduction.

It is known that NaBH$_4$ can undergo hydrolysis in the presence of acid catalysts, releasing H$_2$. This H$_2$ evolution was also observed here in the absence of 4-NP. Interestingly, when using 4-NP the amount of H$_2$ release is significantly diminished to about one-half in accordance with the occurrence of 4-NP reduction simultaneously with the H$_2$ generation. Table S2 summarizes the amount of H$_2$ measured after 5 h of reaction time.

The types and amount of acidic sites in the HKUST-1 parent and QH-240 catalysts were also assessed by temperature-programmed desorption of NH$_3$ (Figure 7 and Table S3). Two NH$_3$ desorption peaks were measured for HKUS at 170 and 303 °C, corresponding to sites of low and medium acid strength, respectively. For QH-240, these peaks appear at 172 and 313 °C. The much higher NH$_3$ adsorption amount measured by TPD for QH-240 as compared to its parent HKUST-1 indicates that the thermal treatment generates an

---

**Figure 6.** UV−vis spectra of 4-NP and NaBH$_4$ solution after 1.0 mg of QH-240 catalyst was added.

**Figure 7.** NH$_3$-TPD analyses of (a) HKUST and (b) QH-240 catalyst.
increase in the total population of Lewis acid sites of about 2.2 times, most of this increase being due to sites of medium strength. This increase in the population and strength of Lewis sites is one of the reasons explaining the higher catalytic activity of QH-240 in the 4-NP reduction. Worth noting is that the pH of the solution under the conditions of Table 1 is about 10. We performed an additional test of QH-240 catalytic activity at pH 8 under conditions otherwise identical to those of Table 1. A 4-NP conversion of 65% was achieved at pH 8, much lower than that at the NaBH4-free pH. This could be due to the reaction of NaBH4 with the H+ used to decrease the pH value, rendering H2 rather than promoting 4-NP reduction. When the pH is lower than 7, the 4-NP solubility is compromised, because 4-nitrophenolate is not formed.

Further characterization of the acid sites was performed by FT-IR spectroscopy monitoring the wavenumber of the characteristic C≡N vibration band of deuterated acetonitrile that is a common test of acidity. The results are presented in Figure 8. As it can be seen there, while HKUST-1 shows a broad and intense band in the 2265 cm⁻¹ region characteristic of physisorbed CD₃CN, QH-240 and QH-260 show a higher-frequency, sharp peak at 2334 cm⁻¹, indicating the presence of Lewis acidity of medium strength. No CD₃CN adsorption was observed for the QH-300 and QH-400 samples corresponding to Cu₃O, which correlates well with their poor catalytic activity. The higher intensity of the 2334 cm⁻¹ peak associated with Lewis acid sites interacting with acetonitrile for QH-260 as compared to QH-240 indicates that, besides acidity, other factors such as surface area and porosity are also playing a role. Note the large decrease in surface area measured from 160 m² g⁻¹ for QH-240 to 5.6 m² g⁻¹ for QH-260 (Table S1). In this way, QH-240 is the sample in which the best compromise among all of the factors has been achieved, resulting in the higher catalytic activity observed for QH-240 in comparison to QH-260 (see Table 1).

Effect of QH-240 Catalyst Dosage. Because the amount of catalyst plays an important role on the reaction rate, the effect of various dosages of QH-240 catalyst on the room-temperature 4-NP reduction was also examined (Table 2 and Figure S17). No change in 4-NP concentration was observed at rt in the absence of catalyst. In addition, the reaction rate of 4-NP reduction increases continuously in the presence of increasing amounts of QH-240. The reduction performance reaches 98.3% within 6 min over 0.2 mg of QH-240. As the dosage of QH-240 catalyst increases up to 1.0 mg, the reduction efficiency reaches 99.8% at a shorter time of 3 min under the same reaction conditions. Afterward, the addition of increasing QH-240 amounts (>1.0 mg) does not result in a significant increase in the reduction rate under the present conditions. Thus, 0.2 mg is chosen as the suitable amount of QH-240 catalyst for subsequent catalytic experiments.

Investigation of the Catalytic Reduction Kinetics. Regarding the reaction kinetics, the key parameters such as the half-life time and the catalytic reaction order were measured both for HKUST and for QH-240 catalysts (Table 3). As expected, when the reaction is carried out in an excess of NaBH4 as reducing agent, the catalytic reduction of 4-NP was found to obey pseudo first-order kinetics using HKUST-1 or QH-240 as catalysts. The plots of ln(C/C₀) versus time for the reduction of 4-NP using QH-240 and HKUST catalysts are provided in Figure S18, 4-NP concentrations higher than 0.05 mM were also tested (Figure 9). The pseudo first-order rate constant values (kₑ) at room temperature for HKUST and QH-240 catalysts were 3.0 × 10⁻⁴ and 1.02 × 10⁻² s⁻¹, respectively. QH-240 catalyst shows a higher activity factor of 51 s⁻¹ g⁻¹ at room temperature. This activity factor is 34 times higher than that of the HKUST catalyst (Table 3). Additionally, the half-life time of the catalytic reduction decreases from 38.5 to 1.13 min using the HKUST and QH-240 catalysts, respectively, which is 34 times faster when using QH-240 catalyst.

The activity difference between the HKUST and QH-240 can be ascribed to the larger quantity of active sites as well as the higher mesopore volume in QH-240, resulting in a more favorable diffusion of reagents to a larger number of catalytic sites. On the other hand, the benzene ring of the ligand in the MOF mesopores can facilitate 4-nitrophenolate adsorption via π−π stacking interaction and hence increase the catalytic

| Table 2. Effect of the QH-240 Amount as a Catalyst on the Reduction Efficiency of 4-NP* |
| catalyst dosage (mg) | 4-NP conversion (%) | reduction time (min)* | 4-AP yield (%) |
|----------------------|---------------------|-----------------------|----------------|
| 1                    | 0                   | 1 week                | 0              |
| 2                    | 0.05                | 75                    | 60             |
| 3                    | 0.1                 | 87                    | 15             |
| 4                    | 0.2                 | 98.3                  | 6              |
| 5                    | 0.5                 | 99.5                  | 4              |
| 6                    | 1.0                 | 99.8                  | 3              |

*Experimental conditions: [4-NP] = 0.05 mM, [NaBH4] = 10 mM, and T = 25 °C. Reduction time = time after subtraction of induction time for each catalytic run. The concentration changes of 4-AP obtained from the absorption changes in λmax = 300 nm.

| Table 3. Comparison of the Apparent Reaction Rate (kapp) and Activity Factor (K) of the HKUST and QH-240 Catalysts for the 4-NP Reduction at Room Temperature |
|----------------------|----------------------|----------------|----------------|
| catalyst             | kapp (s⁻¹)           | t₁/₂ (s)       | K (s⁻¹ g⁻¹)    |
| HKUST                | 3.0 × 10⁻⁴           | 2310           | 1.5            |
| QH-240               | 1.02 × 10⁻²          | 68             | 51             |

ACS Appl. Mater. Interfaces 2022, 14, 978–989
https://doi.org/10.1021/acsami.1c19862
efficiency (Figure S13). Therefore, the synergistic effect of the existence of both meso- and micropores (type I and IV curves) and the greater number of Lewis sites in the catalyst are probably responsible for the better adsorption of 4-NP and BH$_4^-$ ions, reducing the activation kinetic barrier that subsequently results in its superior catalytic performance in comparison with the parent HKUST-1.

**Reaction Mechanism, Rate Constants, and Adsorption Constants of 4-NP and BH$_4^-$ over QH-240.** As was reported in the literature,$^{34,35,41}$ a possible mechanism for the reduction of 4-NP by NaBH$_4$ as reducing agent over a metal catalyst obeys the Langmuir–Hinshelwood (LH) model. In this model, the reaction proceeds having both BH$_4^-$ and nitrophenolate ions adsorbed on the surface of the catalyst, on which the transfer of a hydrogen species would occur. To achieve a complete collection of kinetic data, the apparent rate constant ($k_{app}$) was calculated in a fixed concentration of 4-NP at a given temperature, while varying the NaBH$_4$ concentration. In a second series of kinetic measurements, the concentration of NaBH$_4$ was kept constant, but the 4-NP concentration was varied. Because the temperature is a main parameter in 4-NP reduction, the influence of the reaction temperature was also investigated (Table 4 and Figure 9).

The apparent rate constant is proportional to the surface area of the catalyst ($S$). The kinetic constant can be calculated as

$$k_{app} = \frac{d[NP]}{dt} = -k_{app}c_{NP}$$

For quantitative data analysis, the catalytic reduction can be modeled by the Langmuir–Freundlich isotherm:

$$\theta_i = \frac{(K_i c_i)^n}{1 + \sum_{j=1}^{N} (K_j c_j)}$$

where $\theta_i$ is the surface coverage of the compound $i$, $K_i$ is the adsorption constant of the respective component, $c_i$ is the concentration in solution, and “$n$” is related to the heterogeneity of the sorbent. Further rearrangement of eq 5

Table 4. Rate Constants for 4-NP Reduction and Adsorption Constants of 4-NP and BH$_4^-$ at Different Temperatures on QH-240 Catalyst Determined from the Fitting of the Experimental Data to the LH Model According to Equation 7a

| temp (K) | $k$ [mol/m$^2 \cdot$ s] $\times 10^2$ | $K_{NP}$ [L/mol] | $K_{BH4}$ [L/mol] |
|----------|--------------------------------------|------------------|------------------|
| 298      | 1.0 ± 0.1                            | 2499 ± 130       | 89 ± 11          |
| 303      | 3.0 ± 0.1                            | 3067 ± 175       | 92 ± 14          |
| 308      | 4.6 ± 0.1                            | 3574 ± 250       | 93 ± 13          |
| 313      | 5.8 ± 0.1                            | 4416 ± 265       | 95 ± 15          |

$a$ $n = 0.5$, $m = 1$ Freundlich exponents in eq 7a.
creates eq 7, which can be applied to model the catalytic activity.

\[
-\frac{dc_{\text{NP}}}{dt} = \frac{k_S(K_{\text{NP}}c_{\text{NP}})^n(K_{\text{BH}_4}c_{\text{BH}_4})^m}{(1 + (K_{\text{NP}}c_{\text{NP}})^n + (K_{\text{BH}_4}c_{\text{BH}_4})^m)^2} = k_{\text{app}}c_{\text{NP}}
\]

Thus, \( k_{\text{app}} \) is calculated by

\[
k_{\text{app}} = \frac{k_Sc_{\text{NP}}^n(K_{\text{BH}_4}c_{\text{BH}_4})^m}{(1 + (K_{\text{NP}}c_{\text{NP}})^n + (K_{\text{BH}_4}c_{\text{BH}_4})^m)^2} = c_{\text{NP}}
\]

(7a)

Here, \( k \) is the molar rate constant per square meter of the catalyst, and \( K_{\text{NP}} \) and \( K_{\text{BH}_4} \) are the adsorption coefficients of 4-NP and BH\(_4^-\) respectively.

According to eq 7, an almost complete coverage of the catalyst surface by 4-NP occurs in a high concentration of 4-NP molecules, causing a dramatic decrease in the reaction rate due to the absence of BH\(_4^-\) ions on the active metal sites (Figure 9). In addition, the dependence of \( k_{\text{app}} \) on BH\(_4^-\) concentration is nonlinear (Figure 9), and saturation of \( k_{\text{app}} \) at high BH\(_4^-\) concentrations indicates that there must be a competition of both reactants for the active Cu sites on the catalytic surface. These observations agree with the operation of the LH model for the catalytic 4-NP reduction with QH-240 as catalyst. The rate constant decreases by increasing the concentration of 4-NP and increases by increasing the concentration of NaBH\(_4\), according to their relative adsorption constants. It is obvious that there is a competition between both reactants for active copper sites on the QH-240 surface. The diffusion of the reactants to the unsaturated Cu sites of QH-240 catalyst and the adsorption/desorption equilibrium are surmised to be fast. In the rate-determining step, the surface hydrogen and 4-NP react with each other, and the resulting 4-AP desorbs from the surface of the QH-240 catalyst. Separation of the produced 4-AP releases a free catalytic site, and a new turnover can occur again. A high concentration of 4-NP may cause the full coverage of the catalytic sites, which decreases the reaction rate. Increasing concentrations of NaBH\(_4\) provide an increase in the rate constant until a maximum rate constant is achieved. Beyond this point, an additional NaBH\(_4\) concentration increase decreases the rate constant due to the saturation on the catalytic sites. According to the results, both BH\(_4^-\) ions and 4-NP simultaneously are adsorbed on the surface of the QH-240 catalyst, and then the produced 4-AP is desorbed out to regenerate the active site (Scheme 1). The adsorption constant of 4-NP is higher than that for BH\(_4^-\). Also, \( K_{4\text{-NP}} \) increases upon increasing temperature, while the \( K_{\text{BH}_4^-}\) is comparatively much smaller than the \( K_{4\text{-NP}} \) at each temperature in the range studied (Table 4). Confirmation of the strong 4-NP adsorption on QH-240 was also supported by observation in the IR of blue shifts in the frequency of the characteristic ligand vibrations from 1611 and 1542 to 1614 and 1551 cm\(^{-1}\), respectively, upon adsorption of 4-NP (Figure S13). These adsorption data are compatible with the initial quick reduction of 4-NP to the 4-hydroxylaminophenol intermediate that subsequently is slowly reduced to the final 4-AP product. That means the second step \( (K_{\text{BH}_4^-}) \) is the rate-determining step.

Thermodynamic parameters for the heterogeneous catalytic reduction of 4-NP over QH-240 catalyst were determined by performing the reduction reaction at different temperatures from 25 to 40 °C, obtaining the true \( (k) \) rate constant at each temperature. The true activation energy \( (E_\text{A}) \) was calculated to be 28.8 kJ/mol by the Arrhenius equation:

Scheme 1. Mechanistic Proposal Based on the LH Model for the Reduction of 4-NP by NaBH\(_4^-\) in the Presence of QH-240 Catalyst
\[
\ln(k) = \ln A - \frac{E_a}{RT}
\]  

(8)

With these kinetic data, the activation enthalpy \((\Delta H^\ddagger)\) and activation entropy \((\Delta S^\ddagger)\) were also estimated by the Eyring equation:

\[
\ln \left( \frac{k}{T} \right) = -\frac{\Delta H^\ddagger}{RT} + \frac{\Delta S^\ddagger}{R}
\]

(9)

The Gibbs free energy of activation \(\Delta G^\ddagger\) at 25 °C was also calculated. As presented in Table 5, the adsorption process for the 4-NP molecule is endothermic, while the activation entropy is a positive value, indicating the release of water molecules and/or other adsorbed species bonded to the catalyst surface, which results in an increase of entropy.

The negative \(\Delta G^\ddagger\) value of 4-NP adsorption indicates that the process occurs spontaneously in the presence of this reactant. Also, in the case of NaBH\(_4\) the \(\Delta G^\ddagger\) value is positive, but small, showing that the catalytic conversion is feasible over QH-240 catalyst at room temperature as observed experimentally. The activity factor of the QH-240 catalyst indicates that it is able to contend favorably with diverse reported copper base catalysts (Table 6). Also, the turnover frequency of the QH-240, which is defined as \([(\text{moles of 4-NP reduction})/(\text{moles of active sites}) \times t (\text{min})]\), is at about 0.73 min\(^{-1}\), considering tetrahedral Cu\(^{2+}\) ions as active sites in QH-240.

### Reusability of QH-240 Catalyst.

To examine catalyst reusability, the reduction of 4-NP over QH-240 was carried out for four consecutive runs, calculating the apparent rate constants for each run. The rate constants remain relatively unaltered, even for the fourth run. QH-240 catalyst exhibited a 4-AP yield of up to 98.3% and was recycled with no apparent loss of performance (Figure 10), indicating the stability of the catalyst. Moreover, ICP analysis indicates that the Cu leached from the solid in the aqueous supernatant after each cycle is not observed. These results prove that the QH-240 is stable and can be reused in the catalytic reduction of 4-NP.

### Comparison of the Catalytic Activity of QH-240.

As commented in the Introduction, the purpose of the present study is to determine to which extent the catalytic activity of pristine MOFs, such as HKUST-1 in the present case, can be

---

**Table 5. Activation Parameters for the Adsorption of 4-NP and BH\(_4\) on the Surface of QH-240 Catalyst**

| parameters     | \(K_{4\text{-NP}}\) (kJ/mol) | \(K_{\text{BH}_4}\) (kJ/mol) | \(\Delta H^\ddagger\) (kJ/mol) | \(\Delta S^\ddagger\) (J/mol·K) | \(\Delta G^\ddagger\) (kJ/mol) |
|----------------|-------------------------------|------------------------------|-------------------------------|-------------------------------|-------------------------------|
| \(25.7 \pm 0.2\) | \(1.7 \pm 0.5\)               | \(25.7 \pm 0.2\)             | \(106 \pm 0.6\)              | \(7.8 \pm 0.4\)              | \(−31.6 \pm 0.4\)            |

**Table 6. Comparison of the Activity Factor of QH-240 with Other Reported Copper-Based Catalysts for the Conversion of 4-NP to 4-AP at Room Temperature**

| catalysts                        | \(K_{\text{app}}\) (s\(^{-1}\)) | \(K\) (s\(^{-1}\) g\(^{-1}\)) | \(E\) (kJ/mol) | catalyst dosage (mg) | NaBH\(_4\) (mol)/4-NP (mol) | ref |
|----------------------------------|---------------------------------|---------------------------------|----------------|----------------------|------------------------------|-----|
| Cu\(_x\)O@C-400                  | \(4.8 \times 10^{-3}\)         | 2.4                             | 2              | 2                    | 62.5                         | 42  |
| \((\text{Au}0.3\text{Pt}0.3\text{Pd}0.4)/\text{Cu(HBTC)-1}\) | \(8 \times 10^{-3}\)          | 40                              | 0.2            | 5                    | 25                           | 43  |
| Cu@Cu                           | \(5.9 \times 10^{-2}\)         | 59                              | 1              | 232                  | 44                           |     |
| CuO                             | \(1.3 \times 10^{-3}\)         | 0.26                            | 5              |                      | 45                           |     |
| CuO/Fe\(_3\)O\(_4\)            | \(3.6 \times 10^{-3}\)         | 0.72                            |                |                      |                              |     |
| carbon-doped CuO/Fe\(_3\)O\(_4\) | \(6.5 \times 10^{-3}\)         | 1.3                             |                |                      |                              |     |
| carbonized HKUST/melamine        | \(1.8 \times 10^{-2}\)         | 1.8                             | 73.6           | 0.1                  | 200                          | 46  |
| Ag NPs@ZrGP\(^a\)              | \(1.7 \times 10^{-1}\)         | 17                              | 10             | 100                  | 47                           |     |
| QH-240                          | \(1.02 \times 10^{-2}\)        | 51                              | 28.8           | 0.2                  | 200                          | this study |

\(^a\)ZrGP: zirconium glyphosate.

---
increased by thermal deligandation in quasi-MOFs. The previous reported studies provide relative data among HKUST-1 and the various quasi-MOFs derived therefrom. To put the activity achieved for QH-240 in a broader context, Table 6 compares the reported activity of other catalysts with that achieved herein for QH-240. The reaction under study is suitable for this comparison, because the first-order kinetics allows one to use the apparent first-order rate constant and its specific value activity factor as figures of merit. Although comparisons have always to be taken cautiously due to the possible influence of different factors and conditions, the data shown in Table 6 indicate that QH-240 is among the best material ever reported for this process, thus illustrating the potential of the thermal treatment for the optimization of MOF catalytic activity.

**CONCLUSIONS**

Thermal treatment of HKUST-1 at temperatures between 200 and 300 °C may cause the partial decomposition of the BTC ligands resulting in a “quasi”-MOF material in which the crystal HKUST structure is still partially preserved on the way to CuO. Quasi-HKUST-1 structures act as non-noble hydrogenation catalysts for 4-nitrophenol reduction at room temperature. The higher activity of QH-240 derives from the synergistic effects of the increase in the density of unsaturated copper sites and the existence of both meso- and micropores in the catalyst. The strong adsorption coefficient of 4-NP on QH-240 suggests that phenyl rings of the partially damaged BTC ligands in the MOF mesopores still facilitate 4-NP uptake via π−π stacking, as supported by the blue shifts in the IR vibration frequencies of the benzenetricarboxylate ligands. Thus, the present results show that postsynthetic defective engineering via controlled thermalization of MOFs is a simple and convenient way to further enhance the catalytic activity of MOFs that can be used for the removal of organic pollutants from contaminated waters to convert them into valuable side products.

**ASSOCIATED CONTENT**

Supporting Information

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acsami.1c19862.

FT-IR, Raman, EPR, and UV−vis spectra, SEM images, kinetics plots, and H2 evolution data (PDF)

**AUTHOR INFORMATION**

Corresponding Authors

Mohammad Yaser Masoomi — Department of Chemistry, Faculty of Science, Arak University, Arak 3848177584, Iran; Email: m-masoomi@araku.ac.ir

Hermenegildo Garcia — Instituto Universitario de Tecnología Química Consejo Superior de Investigaciones Científicas and Departamento de Química, Universitat Politècnica de Valencia, Valencia 46022, Spain; orcid.org/0000-0002-9664-493X; Email: hgarcia@upv.es

Authors

Minoo Bagheri — Department of Chemistry, Faculty of Science, Arak University, Arak 3848177584, Iran

Arianna Melillo — Instituto Universitario de Tecnología Química Consejo Superior de Investigaciones Científicas and Departamento de Química, Universitat Politècnica de Valencia, Valencia 46022, Spain

Belen Ferrer — Instituto Universitario de Tecnología Química Consejo Superior de Investigaciones Científicas and Departamento de Química, Universitat Politècnica de Valencia, Valencia 46022, Spain

Complete contact information is available at:

https://pubs.acs.org/10.1021/acsami.1c19862

**Notes**

The authors declare no competing financial interest.

**ACKNOWLEDGMENTS**

This work is based upon research funded by the Iran National Science Foundation (INSF) under project no. 4000089. Also, support of this investigation by Arak University is gratefully acknowledged. The Spanish Ministry of Science and Innovation (Severo Ochoa and RTI2018-98237-C2-1) and Generalitat Valenciana (Prometeo 2021-038) are gratefully acknowledged.

**REFERENCES**

(1) Hira, S. A.; Nallal, M.; Park, K. H. Fabrication of PdAg Nanoparticle Infused Metal-Organic Framework for Electrochemical and Solution-Chemical Reduction and Detection of Toxic 4-Nitrophenol. Sens. Actuators, B, 2019, 298, 126681.

(2) Sahiner, N.; Ozay, H.; Ozay, O.; Aktas, N. A Soft Hydrogel Reactor for Cobalt Nanoparticle Preparation and Use in the Reduction of Nitrophenols. Appl. Catal., B, 2010, 101 (1), 137–143.

(3) Chu, C.; Rao, S.; Ma, Z.; Han, H. Copper and Cobalt Nanoparticles Doped Nitrogen-Containing Carbon Frameworks Derived from Cu-Capsulated ZIF-67 as High-Efficiency Catalyst for Hydrogenation of 4-Nitrophenol. Appl. Catal., B, 2019, 256, 117792.

(4) Sun, J.; Fu, Y.; He, G.; Sun, X.; Wang, X. Catalytic Hydrogenation of Nitrophenols and Nitrotoluenes over a Palladium/Graphene Nanocomposite. Catal. Sci. Technol., 2014, 4 (6), 1742–1748.

(5) Lu, Y.; Mei, Y.; Drechsler, M.; Ballauf, M. Thermosensitive Core−Shell Particles as Carriers for Ag Nanoparticles: Modulating the Catalytic Activity by a Phase Transition in Networks. Angew. Chem., Int. Ed., 2006, 45 (5), 813–816.

(6) Din, M. I.; Khalid, R.; Hussain, Z.; Hussain, T.; Mujahid, A.; Najeeb, I.; Izhav, F. Nanocatalytic Assemblies for Catalytic Reduction of Nitrophenols: A Critical Review. Crit. Rev. Anal. Chem. 2020, 50 (4), 322–338.

(7) Wu, W.; Lei, M.; Yang, S.; Zhou, L.; Liu, L.; Xiao, X.; Jiang, C.; Roy, V. A. L. A One-Pot Route to the Synthesis of Alloyed Cu/Ag Bimetallic Nanoparticles with Different Mass Ratios for Catalytic Reduction of 4-Nitrophenol. J. Mater. Chem. A, 2015, 3 (7), 3450–3455.

(8) Morales, M. V.; Rocha, M.; Freire, C.; Asegedbegha-Nieto, E.; Gallegos-Suarez, E.; Rodriguez-Ramos, I.; Guerrero-Ruiz, A. Development of Highly Efficient Cu Versus Pd Catalysts Supported on Graphitic Carbon Materials for the Reduction of 4-Nitrophenol to 4-Aminophenol at Room Temperature. Carbon, 2017, 111, 150–161.

(9) Liu, W.-J.; Tian, K.; Jiang, H.; Yu, H.-Q. Harvest of Cu Np Anchored Magnetic Carbon Materials from Fe/Cu Preloaded Biomass: Their Pyrolysis, Characterization, and Catalytic Activity on Aqueous Reduction of 4-Nitrophenol. Green Chem. 2014, 16 (9), 4198–4205.

(10) Cai, R.; Ellis, P. R.; Yin, J.; Liu, J.; Brown, C. M.; Griffin, R.; Chang, G.; Yang, D.; Ren, J.; Cooke, K.; Bishop, P. T.; Theis, W.; Palmer, R. E. Performance of Preformed Au/Cu Nanoclusters Deposited on MgO Powders in the Catalytic Reduction of 4-Nitrophenol in Solution. Small, 2018, 14 (13), 1703734.
Organic Frameworks for Catalytic Applications. Coord. Chem. Rev. 2019, 5 (75), 6076–6079.

Gascon, J. Metal with MOFs. Coord. Chem. Rev. 2020, 110 (8), 4606–4655.

Mixed-Metal MOFs: Unique Opportunities in Metal–Organic Framework (MOF) Functionality and Design. Angew. Chem., Int. Ed. 2019, 58 (43), 15188–15205.

Gascon, J. Metal with MOFs. Coord. Chem. Rev. 2020, 113 (5), 1304–1306.

Selective Catalytic Reduction of NOx at Low Temperature. Preparation of Quasi-MIL-101(Cr) Loaded Ceria Catalysts for the Removal Performance of a Cobalt Based Metal-Organic Framework. Catalyst in the Pods Process of Light Oil. Organic Framework Composite as a Simultaneous Photocatalyst and Nanoparticles Immobilized in Spherical Polyelectrolyte Brushes. J. Phys. Chem. C 2010, 114 (19), 8814–8820.

Selective Catalytic Reduction of NOx by Magnetically Recoverable Au Nanocatalyst. J. Hazard. Mater. 2009, 165 (1), 664–669.

Towards Reduction of 4-Nitrophenol. A Multifunctional Tb-MOF for Highly Discriminative Sensing of Eu3+/ Dy3+ and as a Catalyst Support of Ag Nanoparticles. Small 2017, 13 (22), 1602996.

Catalytic Activity of Faceted Gold Nanoparticles Studied by a Model Reaction: Evidence for Substrate-Induced Surface Restructuring. ACS Catal. 2011, 1 (8), 908–916.

One-Pot Green Synthesis of Ag-Decorated SnO2Microsphere: An Efficient and Reusable Catalyst for Reduction of 4-Nitrophenol. Nanoscale Res. Lett. 2017, 12 (1), 435.

Nature of Copper Active Sites in the Carbon Monoxide Oxidation on Cu2Oand Cu2O4spinel Type Catalysts. J. Catal. 1998, 177 (1), 82–95.

Lattic Dynamics of a Prototypical Metal–Organic Framework. Adv. Theory Simul. 2019, 2 (11), 1900093.

Partial Deligandation of M/Ce/Btc Nanorods (M = Au, Cu, Au-Cu) with “Quasi-MOF” Structures Towards Improving Catalytic Activity and Stability. Appl. Catal., A 2019, 572, 34–43.

Nature of Copper Active Sites in the Carbon Monoxide Oxidation on Cu2O and Cu2O4spinel Type Catalysts. J. Catal. 1998, 177 (1), 82–95.

Lattice Dynamics of a Prototypical Metal–Organic Framework. Adv. Theory Simul. 2019, 2 (11), 1900093.

Catalytic Activity of Faceted Gold Nanoparticles Studied by a Model Reaction: Evidence for Substrate-Induced Surface Restructuring. ACS Catal. 2011, 1 (8), 908–916.

Selective Oxidation of Benzyl Alcohol in Air. Materials Chemistry Frontiers 2019, 3 (11), 2363–2373.

Synthesis of Copper Oxide/Reduced Graphene Oxide Nanocomposite and Its Enhanced Catalytic Activity Towards Reduction of 4-Nitrophenol. RSC Adv. 2015, 5 (75), 6076–6079.