Highly Efficient MOF-Driven Silver Subnanometer Clusters for the Catalytic Buchner Ring Expansion Reaction

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ABSTRACT: The preparation of novel efficient catalysts—that could be applicable in industrially important chemical processes—has attracted great interest. Small subnanometer metal clusters can exhibit outstanding catalytic capabilities, and thus, research efforts have been devoted, recently, to synthesize novel catalysts bearing such active sites. Here, we report the gram-scale preparation of Ag$_2$ subnanometer clusters within the channels of a highly crystalline three-dimensional anionic metal–organic framework, with the formula $[\text{Ag}_2\text{mpba}\text{Ni}_{\text{III}}\text{Cu}_{\text{II}}\text{Me}_{\text{mpba}}]_0.5\text{Na}_4\text{H}_{23}\text{O} = \text{Ag}_2\text{Me}_{\text{mpba}}^{1+} = \text{N,N′-2,4,6-trimethyl-1,3-phenylenebis(oxamate)}$. The resulting crystalline solid catalyst—fully characterized with the help of single-crystal X-ray diffraction—exhibits high catalytic activity for the catalytic Buchner ring expansion reaction.

MOFs are crystalline porous materials that have attracted significant attention in the past 2 decades due to the myriad applications they can be used in. Moreover, aiming at encapsulating/synthesizing SNMCs, or even single-atom catalysts, MOFs offer clear advantages compared to other porous materials such as a fine control of the functionalities decorating the channels—which allows us to retain and align metals in specific positions and controlled stoichiometries—and the possibility to use single-crystal X-ray crystallography to unveil the crystal structure of these ultrasmall metal species.

In this context, we recently used, as a chemical reactor, a highly robust anionic three-dimensional MOF, with the formula $\text{Ni}_4\text{Cu}_2\text{Me}_{\text{mpba}}\text{Na}_4\text{H}_{23}\text{O} = \text{N,N′-2,4,6-trimethyl-1,3-phenylenebis(oxamate)}$, for the MOF-driven preparation of ligand-free tetranuclear $\text{Pd}_4^{1+}$ clusters after two consecutive post-synthetic steps consisting of first replacing the Ni$^{2+}$ cations hosted within its channels by Pd$^{2+}$ ones and the concomitant reduction to form the final tetranuclear species within the empty space of the MOF. The resulting host–guest material had the following formula: $[\text{Pd}_4\text{Me}_{\text{mpba}}\text{Na}_4\text{H}_{23}\text{O} = \text{N,N′-2,4,6-trimethyl-1,3-phenylenebis(oxamate)}]$, for the MOF-driven preparation of ligand-free tetranuclear $\text{Pd}_4^{1+}$ clusters after two consecutive post-synthetic steps consisting of first replacing the Ni$^{2+}$ cations hosted within its channels by Pd$^{2+}$ ones and the concomitant reduction to form the final tetranuclear species within the empty space of the MOF.
of the number of inserted Pd²⁺ cations in the first step, which were occupying specific positions by interacting with the carboxylate oxygens from the network. Then, the confined space, as well as the mentioned controlled stoichiometry, allowed the formation of the small [Pd₄]²⁺ clusters, homogeneously distributed within the walls of the MOF (Figure 1). As expected, such naked clusters, possessing all four metal atoms outwardly exposed, exhibited outstanding catalytic activity, outperforming state-of-the-art metal catalysts in carbene-mediated reactions, also showing high yields (>90%) and turnover numbers (up to 100,000). However, considering the high cost of palladium, it would be highly desirable to achieve highly performing SNMCs with lower prices.

**RESULTS AND DISCUSSION**

Herein, aiming at expanding these results to more affordable metals, we report the two-step post-synthetic preparation of Ag₂ nanoclusters using the same MOF Ni²⁺[Ni⁴⁺Cu₆⁺(Me₃mpba)₂]₃·54H₂O (I) as the host matrix (Figure 2a). First, nickel(II) cations, located in the pores of I, are exchanged by Ag⁺ ions, yielding the novel compound Ag₄[I²⁺Cu₈⁺(Me₃mpba)₂]₃·51H₂O (2) (Figure 2b). Then, after introducing NaBH₄, the reduction process occurs to give the final compound [Ag₄]@Ag₂Na₂[Li⁺Cu₆⁺(Me₃mpba)₂]₃·48H₂O (3) (Figure 2c). The whole process could be followed by single-crystal X-ray diffraction (SCXRD), unveiling certain details about the nanocluster formation, which constitutes one of the very few examples of MOF-hosted silver subnanometer clusters and the first whose crystal structure could be elucidated.

The nature of the final hybrid material, 3, containing Ag₂ clusters (together with unreduced Ag⁺ ions) has been further confirmed by the combination of a variety of techniques including inductively coupled plasma—mass spectrometry (ICP—MS) (Table S1, Supporting Information), elemental mapping, powder X-ray diffraction (PXRD), thermogravimetric analysis (TGA), X-ray photoelectron spectroscopy (XPS), and scanning electron microscopy (SEM). The N₂ adsorption isotherms at 77 K confirmed the permanent porosity of 2 and 3 (see below). Finally, as previously mentioned, SCXRD with synchrotron radiation allowed the resolution of the crystal structure of 2 and 3 (Table S2, Supporting Information) and the observation of ultrasmall silver dinuclear entities and surroundings within the solid in 3 (Figures 2 and S1).

The anionic Ni⁴⁺Cu⁴⁺ open-framework structures in both 2 and 3 are isoreticular and crystallize in the P4/nmm space group of the tetragonal system. Compound 2 exhibits the Ag⁺ cations situated within the walls of the hydrophilic octagonal pores (virtual diameter of 2.0 nm), where they are stabilized by noncovalent interactions involving oxamate oxygen atoms [Ag⁺···Oₓanumate of 2.72(1)—2.79(1) Å], with no evidence of previously Ni²⁺ cations of I, thus indicating that they are completely exchanged by Ag⁺ ions (Figure 2b). Ag⁺ ion surroundings unveil interacting oxygen atoms likely belonging to nitrate anions (the whole fragments were not found from the ΔF map, see Supporting Information, Figures S1—S3) or solvent water molecules [Ag⁺···O distance range of 2.38(3)—2.56(3) Å], together with Ag⁺···Ag⁺ with a separation of 2.74(2) Å, which is shorter than the van der Waals contact distance, and they might be considered as precursors of the Ag₂ dimers observed in 3. On the other hand, the crystal structure of 3 reveals the stabilization provided by the nano-confined space of the MOF on the as-synthesized Ag₂ dimers, constricted into the walls of the hydrophilic octagonal channels.
of the anionic Ni\(^{2+}\)-Cu\(^{2+}\) open-framework net (Figures 2c, S4, and S5), together with unreduced Ag\(^{3+}\)–Ag\(^{2+}\) dimers in smaller square pores (Figures 2c and S5). Further hydrated charge-balancing alkali Na\(^+\) cations are retained in the preferential cationic sites, which stabilize the final material, showing an outstanding robustness (Figures S4 and S5). The poorer accessibility to the small square pores for solvated NaBH\(_4\) is most likely the reason for still unreduced Ag\(^{3+}\)–Ag\(^{2+}\) dimers [blocked by Ag\(^{3+}\)–O\(_{\text{oxamate}}\) interactions at a distance of 2.84(1) Å] (Figure S5). Figure 3 shows that Ag\(^{3+}\) dimers

![Image](https://example.com/image1.png)

**Figure 3.** One single channel of 3, showing supramolecular interactions involving oxamate ligands of the network (distances are reported in angstroms).

[intradimer Ag⋯Ag distance of 3.19(1) Å] are well-fixed and stabilized inside the walls of the largest pores of the network by means of supramolecular interactions involving oxamate ligands [Ag⋯O\(_{\text{oxamate}}\) distance range of 2.93(1)–3.05(1) Å] and very weak connections with solvent molecules [Ag\(^{3+}\)⋯O\(_{\text{water}}\) distance of 3.25(1) Å].

SEM coupled with energy-dispersive X-ray spectroscopy (EDX) measurements of 2 and 3 are given in Figures S6 and S7. EDX elemental mappings for Cu, Ni, Ag, and Na (3) elements show a heterogeneous spatial distribution of Ag atoms always located next to Cu and Ni atoms. Moreover, aberration-corrected HAADF–STEM (AC-HAADF–STEM) images are shown in Figure S8. They allow a direct visualization of both Ag\(_2\) dimers together with Ag\(^+\) species—most likely to silver atoms residing in smaller square channels.

TGA of 2 and 3 (Figure S9) established the solvent contents for both materials, which are reflected in their chemical formulas. PXRD patterns of 2 and 3 (Figure S10) indicate that the bulk samples are crystalline and pure, with no typical peaks of Ag\(^{2+}\) nanoparticles. Indeed, experimental diffraction patterns of 2 and 3 are identical to the theoretical ones extracted from the SCXRD data. XPS spectra of compounds 2 and 3 are depicted in Figure S11. For 2, only possessing Ag\(^{2+}\) cations, two bands at 367.6 and 373.6 eV, ascribed to Ag 3d\(_{5/2}\) and Ag 3d\(_{3/2}\) binding energies, respectively, are observed (Figure S11a). In turn, for MOF 3—where SCXRD and elemental analyses suggest that Ag\(^{3+}\) cations and Ag\(^{2+}\) nanoclusters coexist—apart from the same Ag 3d\(_{5/2}\) and Ag 3d\(_{3/2}\) bands at 367.6 and 373.6 eV, respectively, indicative of Ag\(^{2+}\), two additional peaks at 368.4 and 374.4 eV can be observed, which are attributed to reduced Ag\(^0\) atoms, with a 1:1 ratio respect to Ag\(^{2+}\) (Figure S11b). CO-probe diffuse reflectance infrared Fourier transform spectroscopy (DRIFTS) was conducted on MOF 3, run at 77 K to avoid any in situ reduction of Ag\(^{2+}\) and to observe potential Ag\(^0\)–CO species (Figure S12). The results show three main peaks, one at 1938 cm\(^{-1}\), consistent with CO bridged-bonded to Ag\(^0\) atoms,\(^{26}\) a second at 2059 cm\(^{-1}\), attributable to Ag(CO)\(^{+}\) species,\(^{29}\) and a last peak at 2043 cm\(^{-1}\), corresponding to free CO after saturation. It is known that the adsorption of CO on Ag\(^{0}\) is lower than Ag\(^{3+}\);\(^{30}\) thus, the lower intensity of the former makes sense and could very well correspond to a 1:1 ratio between Ag oxidation states. These results strongly support that 50% of Ag\(^{3+}\) present in 2 are reduced by NaBH\(_4\) forming Ag\(^2\) nanoclusters, whereas 50% of Ag\(^{2+}\) cations remain untouched, occupying inaccessible sheltered interstitial positions where the reducing agent cannot access (see the structural description). The N\(_2\) and CO\(_2\) adsorption isotherms for 1–3 confirm their permanent porosity (Figures S13 and S14). N\(_2\) adsorption isotherms for 1–3, with calculated Brunauer–Emmett–Teller\(^{31}\) surface areas of 974, 1013, and 625 m\(^2\)/g, respectively, indicate a very similar permanent porosity for 1 and 3, which is in agreement with their identical estimated virtual diameters of 2.0 nm. In turn, MOF 2 exhibits lower N\(_2\) adsorption despite having the same virtual diameter (2.0 nm), which could be due to a partial collapse of the structure upon solvent evacuation treatment. Remarkably, CO\(_2\) adsorption isotherms show a 66% uptake increase for 3, suggesting quadrupole interactions between CO\(_2\) molecules and Na\(^+\) cations.

The Buchner ring expansion reaction was attempted with catalytic amounts of 3. The results show that the reaction between toluene (4) and ethyl diazoacetate (5) (EDA) proceeds rapidly (30 min) in a very high yield, under standard reaction conditions (Figure 4 top).\(^{31,32}\) Blank experiments without any catalyst gave a 8% conversion, and the use of
MOFs 1 and 2 as catalysts showed the lower catalytic activity of these MOFs than that of 3, with a 3 times lower initial rate for the former (Figure S15). Commercial Ag NPs on alumina only gave a 16% conversion, and remarkably, the state-of-the-art catalyst for this reaction, that is, Rh₃(OAc)₁₄, gave a lower result than MOF 3 in this study, under identical reaction conditions (63% after the addition of 5 at once). An optimum >95% yield of product 6 was obtained after maintaining a low concentration of 5 during the reaction, which was achieved by adding a solution of 5 (in dichloromethane) into the reaction mixture using a syringe pump, instead of adding it at once. Otherwise, the unwanted dimerization reaction of 5 occurs. It is worth noting here that product 6 corresponds to the typical mixture of cycloheptatriene isomers, in accordance with previous results. 24,32 A hot filtration test, where the solid MOF catalyst 3 is removed from the reaction mixture at the reaction temperature (60 °C) at an early conversion (∼30%), shows that the catalytic active species are not present in solution within the experimental error (<10%, Figure 4 bottom), which supports the relative stability of the solid catalyst. In accordance with this result, MOF 3 can be recovered at the end of the reaction by centrifugation, washed, and reused six times, maintaining a good catalytic activity (Figure S16). However, the catalytic yield of MOF 3 decreased to 30% after six uses, which could be due to the progressive (although minor) leaching of active species during the reaction.

Table 1. Results for the Buchner Ring Expansion Reaction between Different Aromatics 7–12 and 5, Catalyzed by MOF 3

| entry | aryl substrate | substituent(s) | product | yield (%) |
|-------|----------------|---------------|---------|-----------|
| 1     | 7              | Cl            | 13      | 66.5      |
| 2     | 8              | Br            | 14      | 64.9      |
| 3     | 9              | CN            | 15      | 92.8      |
| 4     | 10             | OMe           | 16      | 82.6      |
| 5     | 11             | CH₃Br         | 17      | 50.2      |
| 6     | 12             | Me-ortho-F    | 18      | 72.7      |

“GC yields after syringe pump addition of 5.”

Conclusions

In conclusion, ligand-free Ag₃ clusters have been prepared, stabilized, and characterized within an MOF and used as efficient and recoverable catalysts for the Buchner ring expansion reaction. These results expand the toolkit of readily affordable Ag species for heterogeneous catalysis in organic synthesis.

Experimental Section

Preparation of Ag₃(Ni₄(Cu₃(mpba)₃)J-SiH₂O (2). Well-formed deep green prisms of 2, which were suitable for XRD, were obtained by immersing crystals of 1 (ca. 0.0015 mmol) for 48 h in 5 mL of a AgNO₃ aqueous solution (0.004 mmol), which was replaced three times. A multigram scale procedure was also carried out by using the same synthetic procedure but with greater amounts of both, a powder sample of compound 1 (ca. 20 g, 5.8 mmol) and AgNO₃ (2.38 g, 14.0 mmol), with the same successful results and a very high yield (20.33 g, 96%). Anal.: calc. (%) for Cu₄Ni₄Ag₃(C₅H₄N₃)₃F₆ (3707.7): C, 25.27; H, 4.40; N, 4.53. Found: C, 25.34; H, 4.37; N, 4.59 IR (KBr): ν = 3008, 2961 and 2926 cm⁻¹ (C–H), 1601 cm⁻¹ (C=O).

Preparation of [Ag₃Ag₃Na₃(Ni₄(Cu₃(mpba)₃)]−48H₂O (3). Both crystals (ca. 5 mg) and a powder polycrystalline sample of 2 (ca. 10 g) were suspended in 50 mL of a H₂O/CH₃OH (1:2) solution, to which an excess of NaBH₄ was added in 26 fractions (each fraction consisting of 1 mol of NaBH₄ per mole of 2 to give a final NaBH₄/MOF molar ratio of 26 or a NaBH₄/Ag atom molar ratio of 13, which is the same), was added progressively in the space of 72 h. After each addition, the mixture was allowed to react for 1.5 h. After this period, samples were gently washed with a H₂O/CH₃OH solution and filtered on paper, giving high yields (ca. 98%). Anal. Calcld. (%) for Cu₄Ni₄Ag₃Na₃C₅H₄N₃F₆ (3699.6): C, 25.32; H, 4.25; N, 4.54. Found: C, 25.28; H, 4.17; N, 4.59. IR (KBr): ν = 3011, 2971 and 2928 cm⁻¹ (C–H), 1605 cm⁻¹ (C=O).

Gas Adsorption. The N₂ and CO₂ adsorption–desorption isotherms at 77 and 273 K were obtained on polycrystalline samples of 2 and 3 using a BELSORP-mini-X instrument. Samples were first activated with methanol and then evacuated at 348 K during 19 h under 10⁻⁶ Torr prior to their analysis.

Microscopy Measurements. SEM elemental analysis was carried out for 2 and 3 using a HITACHI S-4800 electron microscope coupled with an EDX detector. Data were analyzed using QUANTAX 400.

HAADF–STEM characterization for 3 was performed using an HAADF-FEI-TITAN G2 electron microscope. 5 mg of the material was dispersed in 1 mL of absolute EtOH. Carbon-reinforced copper grids (200 mesh) were submerged into the suspension 30 times and then allowed to dry on air for 24 h.

PXRD Measurements. Polycrystalline samples of 2 and 3 were introduced into 0.5 mm borosilicate capillaries prior to being mounted and aligned on an Empyrean PANalytical powder diffractometer, using Cu Kα radiation (λ = 1.54056 Å). For each sample, five repeated measurements were collected at room temperature (2θ = 2–60°) and merged in a single diffractogram.
**XPS Measurements.** Samples of 2 and 3 were prepared by sticking, without sieving, the samples onto a molybdenum plate using a Scotch tape film, followed by air-drying. Measurements were performed on a K-Alpha XPS system using a monochromatic Al K(α) source (1486.6 eV). As an internal reference for the peak positions in the XPS spectra, the C 1s peak has been set at 284.8 eV.

**DRIFTS of Adsorbed CO.** DRIFTS using CO as a probe molecule was used to evaluate the electronic properties of MOF 3. The experiments have been carried out in a homemade IR cell able to work in the high and low (77 K) temperature ranges. Prior to CO adsorption experiments, the sample was evacuated at 298 K under vacuum (10⁻⁶ mbar) for 1 h. CO adsorption experiments were performed at 77 K in the 0.2–20 mbar range. Spectra were recorded once complete coverage of CO at the specified CO partial pressure was achieved. Deconvolution of the IR spectra has been performed in the Origin software using Gaussian curves where the full width at half-maximum of the individual bands has been taken as a constant. The peak areas are normalized to the sample weight.

**X-ray Crystallographic Data Collection and Structural Refinement.** Crystals of 2 and 3 with ca. 0.06 × 0.08 × 0.08 and 0.08 × 0.12 × 0.12 mm dimensions, respectively, were selected and mounted on a MIFIGEN holder in Paratone oil and very quickly placed on a liquid nitrogen stream cooled at 90 K to avoid the possible degradation upon dehydration. Diffraction data for 2–3 were collected using synchrotron at the I19 beamline of the DIAMOND at λ = 0.6889 Å. Crystallographic details can be found in the Supporting Information.

**General Catalytic Reaction Procedure.** MOF 3 (9.5 mg, 10 mol % Ag) was weighed in a 2 mL vial with a magnetic stirrer, and the aromatic substrate (0.8 mL) was added. Then, the vial was placed in a pre-heated oil bath at 60 °C, and ethyl diazoacetate 5 (0.1 mmol) was added, either at once or using a syringe pump (solution in 0.08 M toluene). The reaction mixture was allowed to react for 0.5 h. The reaction was complete, filtration was carried out to separate the solid catalyst. The mixture was evaporated at 90 °C under vacuum (10⁻³ mbar) for 1 h. CO adsorption experiments were performed in a homemade IR cell able to collect using synchrotron at the I19 beamline of the DIAMOND at λ = 0.6889 Å. Crystallographic details can be found in the Supporting Information.

**ASSOCIATED CONTENT**

**Supporting Information**

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.inorgchem.2c01508.

Experimental preparation, analytical characterization; selected data from SEM/EDX and ICP–MS analyses; summary of crystallographic data of 2 and 3; perspective view along the c crystallographic axis of 2 and 3; details along c and a crystallographic axes of a single octagonal pore in 2; one single channel of 2 showing supramolecular interactions involving oxamate ligands of the network stabilizing Ag⁺ dimers; details along c and a crystallographic axes of a single octagonal pore in 2; backscattered SEM images of 2 and 3 and corresponding EDX mapping for Cu, Ni, and Ag elements; AC-HAADF–STEM image of MOF 3 showing the presence of both Ag single atoms and Ag₂⁺ dimers; TGA curves of 2 and 3 under a dry nitrogen atmosphere; theoretical and experimental PXRD pattern profiles of 2 and 3; XPS spectra of 2 and 3; CO-DRIFTS spectra of MOF 3 with different CO doses; nitrogen sorption and desorption isotherms for the activated compounds 1, 2, 3, and 6; CO₂ sorption and desorption isotherms for the activated compounds 1, 2, and 3; kinetics for the Buchner ring expansion reaction between toluene 4 and ethyl diazoacetate 5 catalyzed by MOF 3; Buchner ring expansion reaction between mesitylene 18 and ethyl diazoacetate 5 catalyzed by either MOF 3 or the Rh(κ₃−OAc)₃ salt; kinetics for the Buchner ring expansion reaction between toluene 4 and ethyl diazoacetate 5 catalyzed by MOF 3 under increasing stirring speeds and the corresponding initial rate–stirring rate correlation; experimental PXRD pattern of 3 after catalytic experiments; and XPS spectra of 3 after catalytic experiments CCDC reference numbers: 2155455 (2) and 2155456 (3) (PDF).

**Accession Codes**

CCDC 2155455–2155456 contain the supplementary crystallographic data for this paper. These data can be obtained free of charge via www.ccdc.cam.ac.uk/data_request/cif, or by emailing data_request@ccdc.cam.ac.uk, or by contacting The Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, UK; fax: +44 1223 336033.

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**Notes**

The authors declare no competing financial interest.

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