

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

Estefanía Tiburcio,^{||} Yongkun Zheng,^{||} Marta Mon,^{*} Nuria Martín,^{*} Jesús Ferrando–Soria, Donatella Armentano, Antonio Leyva–Pérez,^{*} and Emilio Pardo^{*}



Cite This: *Inorg. Chem.* 2022, 61, 11796–11802



Read Online

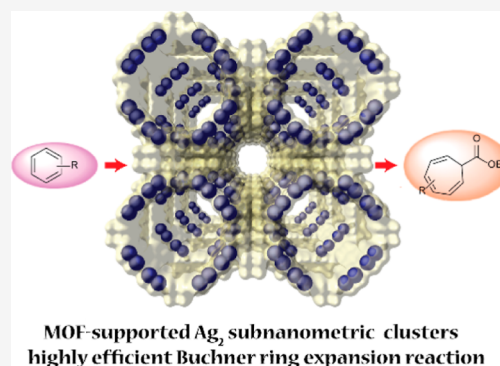
ACCESS |

Metrics & More

Article Recommendations

Supporting Information

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_2^0 subnanometer clusters within the channels of a highly crystalline three-dimensional anionic metal–organic framework, with the formula $[\text{Ag}_2^0]@\text{Ag}_2\text{Na}_2\{\text{Ni}^{\text{II}}_4[\text{Cu}^{\text{II}}_2(\text{Me}_3\text{mpba})_2]_3\}\cdot 48\text{H}_2\text{O}$ [$\text{Me}_3\text{mpba}^{4-} = N,N',2,4,6$ -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.



INTRODUCTION

The preparation, stabilization, and characterization of subnanometer metal clusters (SNMCs) has been a main challenge for chemists during the last years.^{1–7} Indeed, the preparation of such ultrasmall entities is highly complex and often requires, for instance, the use of stabilizing blocking ligands⁸ that prevent their agglomeration into larger metal nanoparticles (MNPs) but, in turn, may worsen their catalytic properties. Moreover, the characterization of such tiny SNMCs is really not an easy task and requires the use of high-resolution microscopy techniques—for example, high-angle annular dark-field–scanning transmission electron microscopy (HAADF–STEM)—whose electron beam can, with time, degrade the sample that is being observed.⁹ However, despite such complications, the outstanding medical, optical, and/or catalytic properties of SNMCs make it well worth the effort.^{3,4,10,11}

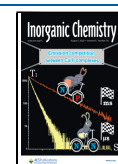
Focusing on the catalytic properties of SNMCs, stabilizing capping ligands of SNMCs dramatically reduces their catalytic activity as they prevent a proper contact with reactants and trigger their decomposition under reaction conditions. Therefore, it seems clear that “naked” SNMCs, with all metal atoms exposed, are called to offer the best catalytic performances.¹² In this context, a proper strategy to obtain ligand-free SNMCs consists in supporting these metal species in porous solids such as zeolites or organic porous materials.¹³ More recently, another type of porous materials, the so-called metal–organic frameworks (MOFs),¹⁴ have emerged as a suitable platform not only to host SNMCs but also to be used as chemical reactors for the in situ chemical synthesis of the SNMCs.^{15,16}

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.¹⁷ Moreover, aiming at encapsulating/synthesizing SNMCs, or even single-atom catalysts,^{18–22} 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^{23–25} to unveil the crystal structure of these ultrasmall metal species.

In this context, we recently used,²¹ as a chemical reactor, a highly robust anionic three-dimensional MOF, with the formula $\text{Ni}^{\text{II}}_2\{\text{Ni}^{\text{II}}_4[\text{Cu}^{\text{II}}_2(\text{Me}_3\text{mpba})_2]_3\}\cdot 54\text{H}_2\text{O}$ [$\text{Me}_3\text{mpba}^{4-} = N,N',2,4,6$ -trimethyl-1,3-phenylenebis(oxamate)], for the MOF-driven preparation of ligand-free tetranuclear $[\text{Pd}_4]^{2+}$ 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: $[\text{Pd}_4]_{0.5}@\text{Na}_3\{\text{Ni}^{\text{II}}_4[\text{Cu}^{\text{II}}_2(\text{Me}_3\text{mpba})_2]_3\}\cdot 56\text{H}_2\text{O}$ (Figure 1). Overall, the anionic nature of the MOF allowed a fine control

Received: May 2, 2022

Published: July 21, 2022



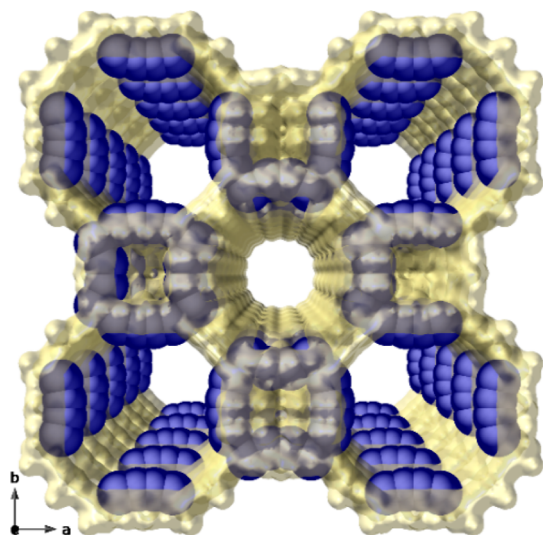


Figure 1. Crystal structure of the previously reported $[\text{Pd}_4]_{0.5}@\text{Na}_3\{\text{Ni}^{\text{II}}_4[\text{Cu}^{\text{II}}_2(\text{Me}_3\text{mpba})_2]_3\}\cdot 56\text{H}_2\text{O}$ MOF.²¹ Atoms constituting the framework and Pd atoms are represented by light yellow and blue spheres, respectively.

of the number of inserted Pd^{2+} 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 $[\text{Pd}_4]^{2+}$ 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 Ag_2^0 nanoclusters using the same MOF $\text{Ni}^{\text{II}}_2\{\text{Ni}^{\text{II}}_4[\text{Cu}^{\text{II}}_2(\text{Me}_3\text{mpba})_2]_3\}\cdot 54\text{H}_2\text{O}$ (1) as the host matrix (Figure 2a). First, nickel(II) cations, located in the pores of 1,

are exchanged by Ag^+ ones, yielding the novel compound $\text{Ag}^+_4\{\text{Ni}^{\text{II}}_4[\text{Cu}^{\text{II}}_2(\text{Me}_3\text{mpba})_2]_3\}\cdot 51\text{H}_2\text{O}$ (2) (Figure 2b). Then, after introducing NaBH_4 , the reduction process occurs to give the final compound $[\text{Ag}_2^0]@\text{Ag}^+_2\text{Na}^+_2\{\text{Ni}^{\text{II}}_4[\text{Cu}^{\text{II}}_2(\text{Me}_3\text{mpba})_2]_3\}\cdot 48\text{H}_2\text{O}$ (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 clusters²⁶ and the first whose crystal structure could be elucidated.

The nature of the final hybrid material, 3, containing Ag_2 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 N_2 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 $\text{Ni}^{\text{II}}_4\text{Cu}^{\text{II}}_6$ open-framework structures in both 2 and 3 are isorecticular and crystallize in the $P4/mmm$ 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 [$\text{Ag}^+\cdots\text{O}_{\text{oxamate}}$ of 2.72(1)–2.79(1) Å], with no evidence of previously Ni^{2+} cations of 1, thus indicating that they are completely exchanged by Ag^+ ones (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 [$\text{Ag}^+\cdots\text{O}$ distance range of 2.38(3)–2.56(3) Å], together with $\text{Ag}^+\cdots\text{Ag}^+$ 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 Ag_2^0 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 Ag_2^0 dimers, constricted into the walls of the hydrophilic octagonal channels

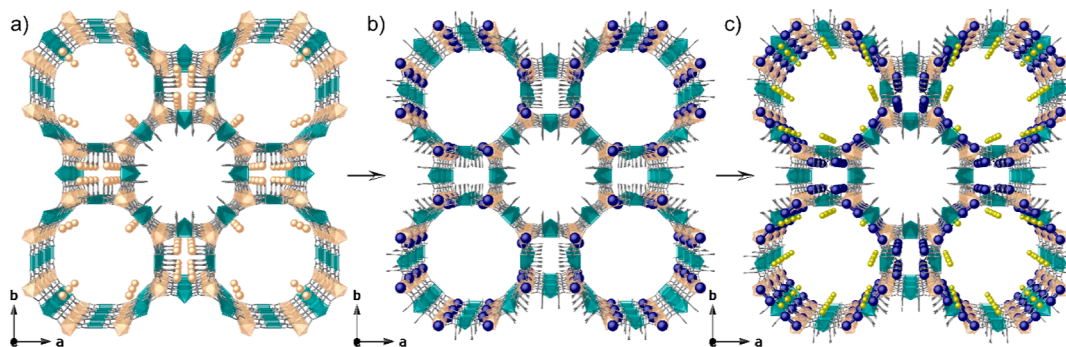


Figure 2. Design approach showing the crystal structures of 1 (a), 2 (b), and 3 (c) showing the two-step post-synthetic process consisting of the exchange of the Ni^{II} cations in the pores of 1 by Ag^+ ones to yield 2 and the reduction process to form the Ag_2^0 clusters in 3. Virtual diameters of larger octagonal pores are 2.0 nm for MOFs 1–3, respectively. Copper and nickel atoms from the network are represented by cyan and orange polyhedra, respectively, whereas organic ligands are depicted as gray sticks. Orange, yellow, and blue spheres represent Ni, Na, and Ag atoms, respectively.

of the anionic $\text{Ni}_4^{2+}\text{Cu}_6^{2+}$ open-framework net (Figures 2c, S4, and S5), together with unreduced $\text{Ag}^+\cdots\text{Ag}^+$ dimers in smaller square pores (Figures 2c and S5). Further hydrated charge-counterbalancing 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 $\text{Ag}^+\cdots\text{Ag}^+$ dimers [blocked by $\text{Ag}^+\cdots\text{O}_{\text{oxamate}}$ interactions at a distance of $2.84(1)$ Å] (Figure S5). Figure 3 shows that Ag_2^0 dimers

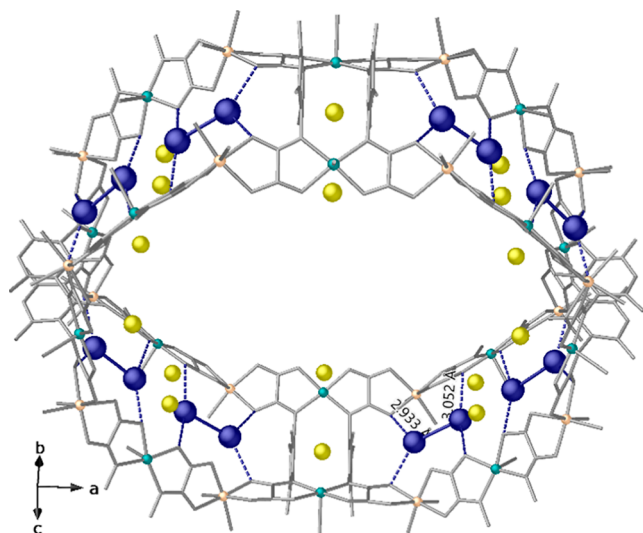


Figure 3. One single channel of 3, showing supramolecular interactions involving oxamate ligands of the network (distances are reported in angstroms).

[intradimer $\text{Ag}\cdots\text{Ag}$ 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 [$\text{Ag}\cdots\text{O}_{\text{oxamate}}$ distance range of $2.93(1)$ – $3.05(1)$ Å] and very weak connections with solvent molecules [$\text{Ag}^0\cdots\text{O}_{\text{water}}$ 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 Ag₁ 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^0 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^+ cations, two bands at 367.6 and 373.6 eV, ascribed to $\text{Ag } 3d_{5/2}$ and $\text{Ag } 3d_{3/2}$ binding energies, respectively,²⁷ are observed (Figure S11a). In turn, for MOF 3—where SCXRD and elemental analyses suggest that Ag^+ cations and Ag_2^0 nanoclusters coexist—apart from the same $\text{Ag } 3d_{5/2}$ and $\text{Ag } 3d_{3/2}$ bands at 367.6 and 373.6 eV, respectively, indicative of Ag^+ , 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^+ (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^+ 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,²⁸ a second at 2059 cm^{-1} , attributable to $\text{Ag}(\text{CO})^+$ 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^+ ;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^+ present in 2 are reduced by NaBH_4 forming Ag_2^0 nanoclusters, whereas 50% of Ag^+ cations remain untouched, occupying inaccessible sheltered interstitial positions where the reducing agent cannot accede (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–Teller³¹ surface areas of 974, 1013, and $625\text{ m}^2/\text{g}$, 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).^{21,32} Blank experiments without any catalyst gave a 8% conversion, and the use of

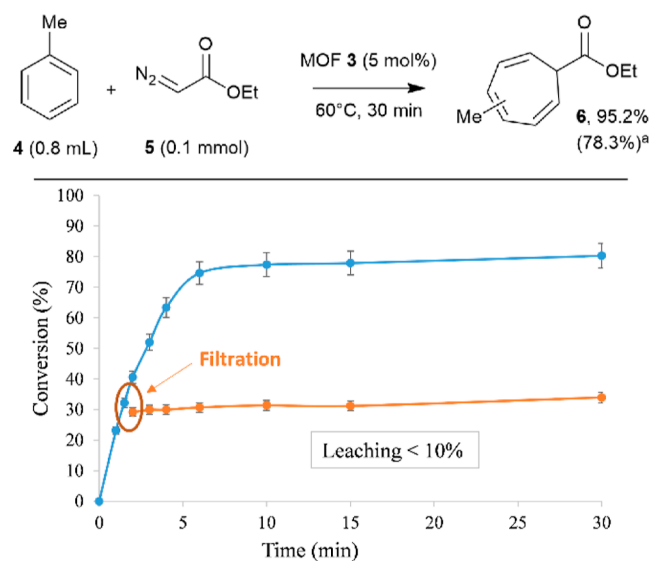
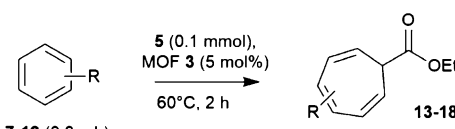


Figure 4. Top: results for the Buchner ring expansion reaction between toluene 4 and ethyl diazoacetate 5 catalyzed by MOF 3. Bottom: Hot filtration test after adding 5 at once, error bars account for 5% uncertainty. (a) GC yield after syringe pump addition of 5, with the result after the addition of 5 at once given in parentheses; the result with $\text{Rh}_2(\text{OAc})_4$ as a catalyst (5 mol %) is 63%.

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, Rh₂(OAc)₄, 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.^{21,32} 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.

The Buchner ring expansion reaction catalyzed by **3** could be expanded to other aromatic substrates (Table 1).

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



entry	aryl substrate	substituent(s)	product	yield (%) ^a
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	CH ₂ Br	17	50.2
6	12	Me-ortho-F	18	72.7

^aGC yields after syringe pump addition of **5**.

Halogenated (products **13**, **14**, and **17**), cyano (product **15**), methoxy (product **16**), and ortho-disubstituted (product **18**) aromatic compounds react with **5** in good to excellent yields and still in short reaction times (<2 h). These results should be remarked upon as this is difficult to find in the open literature Ag-catalyzed Buchner ring expansion reactions.^{32–35} The slightly lower results obtained for the bigger substrates **11** and **12** can be explained by the size discrimination associated to the microporous structure of MOF **3**. To check this, 1,3,5-triisopropylbenzene **19** was tested as a substrate for the reaction, and product **20** was not found (Figure S17). Besides, diffusion tests with varying stirring speeds confirmed that the initial reaction rate is dependent on the stirring, which confirms that the reaction occurs inside the MOF's channels (Figure S18). Indeed, 1,3,5-triisopropylbenzene **19** reacted

when the Rh₂(OAc)₄ salt was used as a catalyst (3 mol %, 69% yield, Figure S17). Therefore, for MOF- or zeolite-stabilized Pd clusters,^{21,36} the formation of Ag₂ clusters in microstructured solids enables not only the catalytic use of this metal in this complex organic synthetic reaction but also its recovery and reuse.^{36,37} Interstitial Ag⁺ cations must be excluded as catalytically active species since reagents cannot access these sites; besides, leaching does not occur under the reaction conditions employed. Finally, the integrity of MOF **3** is ensured by PXRD (Figure S19) and XPS (Figure S20) after catalytic experiments.

CONCLUSIONS

In conclusion, ligand-free Ag₂⁰ 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 Ag₂{Ni^{II}₄[Cu^{II}(Me₃mpba)₂]₃}·51H₂O (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.: calcd (%) for Cu₆Ni₄Ag₄C₇₈H₁₆₂N₁₂O₈₇ (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 [Ag₂]Ag₂Na₂{Ni^{II}₄[Cu^{II}(Me₃mpba)₂]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, divided 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. Calcd (%) for Cu₆Ni₄Ag₄Na₂C₇₈H₁₅₆N₁₂O₈₄ (3699.61): 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 re-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. $0.06 \times 0.08 \times 0.08$ and $0.08 \times 0.12 \times 0.12$ mm dimensions, respectively, were selected and mounted on a MITIGEN 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 $\lambda = 0.6889$ 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 dichloromethane). The mixture was allowed to react for 0.5–2 h. After the reaction was complete, filtration was carried out to separate the solid catalyst. The reaction mixture was analyzed by gas chromatography (GC) and GC–MS. Further details can be found in the [Supporting Information](#).

■ ASSOCIATED CONTENT

SI 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**, and **3**; 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 MOFs **1** and **3**; catalytic reuses 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 Rh₂(OAc)₄ 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.

■ AUTHOR INFORMATION

Corresponding Authors

Marta Mon – Instituto de Tecnología Química (UPV–CSIC), Universidad Politécnica de Valencia–Consejo Superior de Investigaciones Científicas, Valencia 46022, Spain; Email: marmoco@itq.upv.es

Nuria Martín – Departamento de Química Inorgánica, Instituto de Ciencia Molecular (ICMOL), Universidad de Valencia, Valencia 46980, Spain; Email: nuria.martin@uv.es

Antonio Leyva–Pérez – Instituto de Tecnología Química (UPV–CSIC), Universidad Politécnica de Valencia–Consejo Superior de Investigaciones Científicas, Valencia 46022, Spain; orcid.org/0000-0003-1063-5811; Email: anleyva@itq.upv.es

Emilio Pardo – Departamento de Química Inorgánica, Instituto de Ciencia Molecular (ICMOL), Universidad de Valencia, Valencia 46980, Spain; orcid.org/0000-0002-1394-2553; Email: emilio.pardo@uv.es

Authors

Estefanía Tiburcio – Departamento de Química Inorgánica, Instituto de Ciencia Molecular (ICMOL), Universidad de Valencia, Valencia 46980, Spain

Yongkun Zheng – Instituto de Tecnología Química (UPV–CSIC), Universidad Politécnica de Valencia–Consejo Superior de Investigaciones Científicas, Valencia 46022, Spain

Jesús Ferrando–Soria – Departamento de Química Inorgánica, Instituto de Ciencia Molecular (ICMOL), Universidad de Valencia, Valencia 46980, Spain

Donatella Armentano – Dipartimento di Chimica e Tecnologie Chimiche, Università della Calabria, Rende, Cosenza 87036, Italy; orcid.org/0000-0002-8502-8074

Complete contact information is available at:

<https://pubs.acs.org/doi/10.1021/acs.inorgchem.2c01508>

Author Contributions

^{||}E.T. and Y.Z. are contributed equally to this work.

Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

This work was supported by the Ministero dell'Università e della Ricerca (Italy), the MICIIN (Spain) (projects PID2020-

115100GB-I00, PID2019-104778GB-I00 and Excellence Unit “Maria de Maeztu” CEX2019-000919-M), and the Generalitat Valenciana (project PROMETEO/2021/054). Thanks are also extended to the Ramon y Cajal Program (RYC2019-027940-I) (J. F.-S.) and the Diamond Light Source for the awarded beamtime and provision of synchrotron radiation facilities (proposal no. CY22411-1). We particularly acknowledge Dr David Allan and Sarah Barnett for their assistance at the I19 beamline. E.P. acknowledges the financial support of the European Research Council under the European Union’s Horizon 2020 research and innovation programme/ERC grant agreement no 814804, MOF-reactors. M.M. thanks MICIIN from a contract under the Juan de la Cierva program (FJC2019-040523-I). Y.Z. thanks the China Scholarship Council (CSC no: 202009350009) for a Ph.D. fellowship.

REFERENCES

- (1) Liu, L.; Corma, A. Metal Catalysts for Heterogeneous Catalysis: From Single Atoms to Nanoclusters and Nanoparticles. *Chem. Rev.* **2018**, *118*, 4981–5079.
- (2) Li, Z.; Ji, S.; Liu, Y.; Cao, X.; Tian, S.; Chen, Y.; Niu, Z.; Li, Y. Well-Defined Materials for Heterogeneous Catalysis: From Nanoparticles to Isolated Single-Atom Sites. *Chem. Rev.* **2020**, *120*, 623–682.
- (3) Mitchell, S.; Qin, R.; Zheng, N.; Pérez-Ramírez, J. Nanoscale Engineering of Catalytic Materials for Sustainable Technologies. *Nat. Nanotechnol.* **2021**, *16*, 129–139.
- (4) Wang, N.; Sun, Q.; Yu, J. Ultrasmall Metal Nanoparticles Confined within Crystalline Nanoporous Materials: A Fascinating Class of Nanocatalysts. *Adv. Mater.* **2019**, *31*, 1803966.
- (5) Xu, L.; Ma, W.; Wang, L.; Xu, C.; Kuang, H.; Kotov, N. A. Nanoparticle Assemblies: Dimensional Transformation of Nanomaterials and Scalability. *Chem. Soc. Rev.* **2013**, *42*, 3114.
- (6) Grzelczak, M.; Pérez-Juste, J.; Mulvaney, P.; Liz-Marzán, L. M. Shape Control in Gold Nanoparticle Synthesis. *Chem. Soc. Rev.* **2008**, *37*, 1783.
- (7) Perezjuste, J.; Pastorizasantos, I.; Lizmarzan, L.; Mulvaney, P. Gold Nanorods: Synthesis, Characterization and Applications. *Coord. Chem. Rev.* **2005**, *249*, 1870–1901.
- (8) Kang, X.; Wang, S.; Song, Y.; Jin, S.; Sun, G.; Yu, H.; Zhu, M. Bimetallic Au₂Cu₆ Nanoclusters: Strong Luminescence Induced by the Aggregation of Copper(I) Complexes with Gold(0) Species. *Angew. Chem., Int. Ed.* **2016**, *55*, 3611–3614.
- (9) Egerton, R. F.; Li, P.; Malac, M. Radiation Damage in the TEM and SEM. *Micron* **2004**, *35*, 399–409.
- (10) Ni, B.; Wang, X. Chemistry and Properties at a Sub-Nanometer Scale. *Chem. Sci.* **2016**, *7*, 3978–3991.
- (11) Shang, L.; Dong, S.; Nienhaus, G. U. Ultra-Small Fluorescent Metal Nanoclusters: Synthesis and Biological Applications. *Nano Today* **2011**, *6*, 401–418.
- (12) Boronat, M.; Leyva-Pérez, A.; Corma, A. Theoretical and Experimental Insights into the Origin of the Catalytic Activity of Subnanometric Gold Clusters: Attempts to Predict Reactivity with Clusters and Nanoparticles of Gold. *Acc. Chem. Res.* **2014**, *47*, 834–844.
- (13) Liu, L.; Lopez-Haro, M.; Lopes, C. W.; Li, C.; Concepcion, P.; Simonelli, L.; Calvino, J. J.; Corma, A. Regioselective Generation and Reactivity Control of Subnanometric Platinum Clusters in Zeolites for High-Temperature Catalysis. *Nat. Mater.* **2019**, *18*, 866–873.
- (14) Furukawa, H.; Cordova, K. E.; O’Keeffe, M.; Yaghi, O. M. The Chemistry and Applications of Metal–Organic Frameworks. *Science* **2013**, *341*, 974.
- (15) Viciano-Chumillas, M.; Mon, M.; Ferrando-Soria, J.; Corma, A.; Leyva-Pérez, A.; Armentano, D.; Pardo, E. Metal–Organic Frameworks as Chemical Nanoreactors: Synthesis and Stabilization of Catalytically Active Metal Species in Confined Spaces. *Acc. Chem. Res.* **2020**, *53*, 520–531.
- (16) Young, R. J.; Huxley, M. T.; Pardo, E.; Champness, N. R.; Sumbly, C. J.; Doonan, C. J. Isolating Reactive Metal-Based Species in Metal–Organic Frameworks – Viable Strategies and Opportunities. *Chem. Sci.* **2020**, *11*, 4031–4050.
- (17) Freund, R.; Zaremba, O.; Arnauts, G.; Ameloot, R.; Skorupskii, G.; Dincă, M.; Bavykina, A.; Gascon, J.; Ejsmont, A.; Goscianska, J.; Kalmutzki, M.; Lächelt, U.; Ploetz, E.; Diercks, C. S.; Wuttke, S. The Current Status of MOF and COF Applications. *Angew. Chem., Int. Ed.* **2021**, *60*, 23975–24001.
- (18) Bilanin, C.; Tiburcio, E.; Ferrando-Soria, J.; Armentano, D.; Leyva-Pérez, A.; Pardo, E. Crystallographic Visualization of a Double Water Molecule Addition on a Pt¹⁺-MOF during the Low-temperature Water-Gas Shift Reaction. *ChemCatChem* **2021**, *13*, 1195–1200.
- (19) Tiburcio, E.; Greco, R.; Mon, M.; Ballesteros-Soberanas, J.; Ferrando-Soria, J.; López-Haro, M.; Hernández-Garrido, J. C.; Oliver-Meseguer, J.; Marini, C.; Boronat, M.; Armentano, D.; Leyva-Pérez, A.; Pardo, E. Soluble/MOF-Supported Palladium Single Atoms Catalyze the Ligand-, Additive-, and Solvent-Free Aerobic Oxidation of Benzyl Alcohols to Benzoic Acids. *J. Am. Chem. Soc.* **2021**, *143*, 2581–2592.
- (20) Rivero-Crespo, M. A.; Mon, M.; Ferrando-Soria, J.; Lopes, C. W.; Boronat, M.; Leyva-Pérez, A.; Corma, A.; Hernández-Garrido, J. C.; López-Haro, M.; Calvino, J. J.; Ramos-Fernandez, E. V.; Armentano, D.; Pardo, E. Confined Pt¹⁺ Water Clusters in a MOF Catalyze the Low-Temperature Water–Gas Shift Reaction with Both CO₂ Oxygen Atoms Coming from Water. *Angew. Chem., Int. Ed.* **2018**, *57*, 17094–17099.
- (21) Fortea-Pérez, F. R.; Mon, M.; Ferrando-Soria, J.; Boronat, M.; Leyva-Pérez, A.; Corma, A.; Herrera, J. M.; Osadchii, D.; Gascon, J.; Armentano, D.; Pardo, E. The MOF-Driven Synthesis of Supported Palladium Clusters with Catalytic Activity for Carbene-Mediated Chemistry. *Nat. Mater.* **2017**, *16*, 760–766.
- (22) Mon, M.; Rivero-Crespo, M. A.; Ferrando-Soria, J.; Vidal-Moya, A.; Boronat, M.; Leyva-Pérez, A.; Corma, A.; Hernández-Garrido, J. C.; López-Haro, M.; Calvino, J. J.; Ragazzon, G.; Credi, A.; Armentano, A.; Pardo, E. Synthesis of Densely Packaged, Ultrasmall Pt⁰ Clusters within a Thioether-Functionalized MOF: Catalytic Activity in Industrial Reactions at Low Temperature. *Angew. Chem., Int. Ed.* **2018**, *57*, 6186–6191.
- (23) Burgun, A.; Coghlan, C. J.; Huang, D. M.; Chen, W.; Horike, S.; Kitagawa, S.; Alvino, J. F.; Metha, G. F.; Sumbly, C. J.; Doonan, C. J. Mapping-Out Catalytic Processes in a Metal–Organic Framework with Single-Crystal X-Ray Crystallography. *Angew. Chem., Int. Ed.* **2017**, *56*, 8412–8416.
- (24) Inokuma, Y.; Yoshioka, S.; Ariyoshi, J.; Arai, T.; Hitora, Y.; Takada, K.; Matsunaga, S.; Rissanen, K.; Fujita, M. X-Ray Analysis on the Nanogram to Microgram Scale Using Porous Complexes. *Nature* **2013**, *495*, 461–466.
- (25) Mon, M.; Bruno, R.; Ferrando-Soria, J.; Bartella, L.; Di Donna, L.; Talia, M.; Lappano, R.; Maggolini, M.; Armentano, D.; Pardo, E. Crystallographic Snapshots of Host–Guest Interactions in Drugs@metal–Organic Frameworks: Towards Mimicking Molecular Recognition Processes. *Mater. Horiz.* **2018**, *5*, 683–690.
- (26) Zhuang, P.; Zhang, P.; Li, K.; Kumari, B.; Li, D.; Mei, X. Silver Nanoclusters Encapsulated into Metal–Organic Frameworks for Rapid Removal of Heavy Metal Ions from Water. *Molecules* **2019**, *24*, 2442.
- (27) Ferraria, A. M.; Carapeto, A. P.; do Rego, A. M. B. X-Ray Photoelectron Spectroscopy: Silver Salts Revisited. *Vacuum* **2012**, *86*, 1988–1991.
- (28) Müslehiddinoglu, J. CO Adsorption on Supported and Promoted Ag Epoxidation Catalysts. *J. Catal.* **2003**, *213*, 305–320.
- (29) Tarach, K.; Góra-Marek, K.; Chrzan, M.; Walas, S. Quantification of Silver Sites in Zeolites: Carbon Monoxide Sorption Monitored by IR Spectroscopy. *J. Phys. Chem. C* **2014**, *118*, 23751–23760.
- (30) Bartolomeu, R.; Azambre, B.; Westermann, A.; Fernandes, A.; Bértolo, R.; Hamoud, H. I.; Henriques, C.; Da Costa, P.; Ribeiro, F.

Investigation of the Nature of Silver Species on Different Ag-Containing NO_x Reduction Catalysts: On the Effect of the Support. *Appl. Catal., B* **2014**, *150-151*, 204–217.

(31) De Lange, M. F.; Vlugt, T. J. H.; Gascon, J.; Kapteijn, F. Adsorptive Characterization of Porous Solids: Error Analysis Guides the Way. *Microporous Mesoporous Mater.* **2014**, *200*, 199–215.

(32) Díaz-Requejo, M. M.; Pérez, P. J. Copper, Silver and Gold-Based Catalysts for Carbene Addition or Insertion Reactions. *J. Organomet. Chem.* **2005**, *690*, 5441–5450.

(33) Lovely, C. J.; Browning, R. G.; Badarinarayana, V.; Dias, H. V. R. A Silver-Catalyzed Büchner Reaction. *Tetrahedron Lett.* **2005**, *46*, 2453–2455.

(34) Deng, Y.; Jing, C.; Zavalij, P. Y.; Doyle, M. P. Hg(OTf)₂ Catalyzed Intramolecular 1,4-Addition of Donor–Acceptor Cyclopropenes to Arenes. *Org. Lett.* **2015**, *17*, 4312–4315.

(35) Radhika, S.; Abdulla, C. M. A.; Aneja, T.; Anilkumar, G. Silver-Catalysed C–H Bond Activation: A Recent Review. *New J. Chem.* **2021**, *45*, 15718–15738.

(36) Garnes-Portolés, F.; Greco, R.; Oliver-Meseguer, J.; Castellanos-Soriano, J.; Consuelo Jiménez, M.; López-Haro, M.; Hernández-Garrido, J. C.; Boronat, M.; Pérez-Ruiz, R.; Leyva-Pérez, A. Regioirregular and catalytic Mizoroki-Heck reactions. *Nat. Catal.* **2021**, *4*, 293–303.

(37) Oliver-Meseguer, J.; Boronat, M.; Vidal-Moya, A.; Concepción, P.; Rivero-Crespo, M. A.; Leyva-Pérez, A.; Corma, A. Generation and Reactivity of Electron-Rich Carbenes on the Surface of Catalytic Gold Nanoparticles. *J. Am. Chem. Soc.* **2018**, *140*, 3215–3218.