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Single crystalline hollow metal–organic frameworks: a metal–organic polyhedron single crystal as a sacrificial template†

Hyehyun Kim,^a Minhak Oh,^a Dongwook Kim,^a Jeongin Park,^a Junmo Seong,^a
Sang Kyu Kwak^b and Myoung Soo Lah^{*a}

Single crystalline hollow metal–organic frameworks (MOFs) with cavity dimensions on the order of several micrometers and hundreds of micrometers were prepared using a metal–organic polyhedron single crystal as a sacrificial hard template. The hollow nature of the MOF crystal was confirmed by scanning electron microscopy of the crystal sliced using a focused ion beam.

Hollow structures with cavities of nano-/micrometer dimensions at the center of the particles are of great importance because of their fascinating characteristics such as low density, high surface-to-volume ratio, low thermal expansion coefficient and refractive index, and high loading capacity.^{1–3} So far, various fabrication methods have been developed to synthesize the hollow metal/metal oxide nano-/microstructures.⁴

Metal–organic frameworks (MOFs) have received enormous attention from researchers during the past couple of decades as a new class of porous materials, not only because of their high surface areas, but also because of their diversity and tunability.^{5–7} Porous MOFs are usually crystalline materials and their pore dimensions can be modulated by the length of their organic building units.^{8,9} However, it remains extremely difficult to synthesize MOFs with pore dimensions much larger than several nanometers. The largest pore dimension for MOFs reported to date is about 10 nm for IRMOF-74-XI.¹⁰

Several hollow MOF structures have also been prepared *via* synthetic strategies similar to those employed for the preparation of the hollow metal/metal oxide nano-/microstructures. Hollow MOF microspheres can be prepared using surface-modified polystyrene

(PS) microspheres as a hard solid template,^{11,12} emulsion droplets of PS membrane¹³ and Tween-85¹⁴ as soft liquid templates, and CO₂ bubbles¹⁵ as a soft gas template. The droplets of aqueous metal solution injected into the flowing ligand-containing organic solution serve as a sacrificial template and hollow MOF capsules can be produced at the interface of the aqueous droplet and the organic solution.¹⁶ As inorganic hollow nano-/microstructures, hollow MOF nano-/microspheres can be obtained *via* template-free approaches such as Ostwald ripening¹⁷ and a spray-drying strategy.¹⁸ Not all of the reported hollow MOF nano-/microstructures are single crystalline materials. The shell of the hollow MOFs is either an aggregate of smaller non-hollow MOF crystals or a composite of the polymer and the smaller non-hollow MOF crystals. Selective chemical etching of the core of a surface-protected MOF single crystal is a template-free approach to a hollow MOF structure.¹⁹ However, the hollow MOF structure from a single crystalline MOF template is not a single crystalline hollow MOF because the shell of a hollow MOF will be significantly deteriorated during the etching process. Very recently, a hollow MOF with well-defined surface morphology was prepared using a solvothermal reaction without a template and the formation of a hollow cage was proposed through a surface-energy-driven mechanism.²⁰ However, the identity of the hollow MOF proposed as Fe-substituted MOF-5 is not certain. While the molar ratio of Fe to Zn in the proposed MOF is ~3.4, it is well known that the maximum molar ratio of M to Zn in M-substituted MOF-5 is ~0.33.²¹

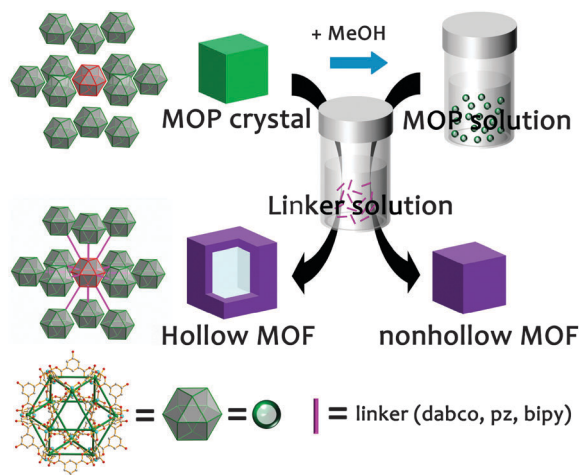
Here, we report for the first time the preparation of single crystalline hollow MOFs with a shell pore of nanometer dimension and hollow cavity of micrometer dimension *via* a stepwise synthetic approach using a metal–organic polyhedron (MOP) single crystal as a hard sacrificial template (Scheme 1). First, we prepared cuboctahedral Cu–MOP single crystals with dimensions of either several micrometers or several hundred micrometers made of approximately 3 nm MOP building units, [Cu₂₄(hip)₂₄S₂₄] (where hip = 5-hydroxyisophthalate). Single MOP crystals were used as hard sacrificial templates. The cuboctahedral Cu–MOP building units of the MOP single crystal as one type of reactant could be interlinked

^a Department of Chemistry, Ulsan National Institute of Science and Technology, Ulsan, 689-798, Korea. E-mail: mslah@unist.ac.kr; Fax: +82 52 217 2019; Tel: +82 52 217 2931

^b Department of Chemical Engineering, Ulsan National Institute of Science and Technology, Ulsan, 689-798, Korea

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Scheme 1 A MOP single crystal as a sacrificial template for a hollow MOF single crystal.

using linear ditopic linkers (1,4-diazabicyclo[2.2.2]octane (dabco), pyrazine (pz), and 4,4'-bipyridine (bipy)) as another type of reactant.

The cuboctahedral Cu-MOP crystals, **MOP-micro** and **MOP-macro**, are obtained from the reactions of Cu(II) ions with H_2hip as a bent ditopic dicarboxylate ligand in DMF (Fig. S1–S5, ESI†). While a solvothermal reaction without stirring resulted in single crystals of several hundred micrometers of **MOP-macro**, a similar reaction at ambient temperature with stirring produced single crystals of **MOP-micro** with dimensions of several micrometers.

Single crystalline hollow MOFs of several hundred micrometer dimensions could be prepared by reacting **MOP-macro** crystals of several hundred micrometer dimensions with ditopic linkers in MeOH, where the **MOP-macro** crystal serves as a sacrificial template. The MOP crystal serves both as a crystalline template and as a reactant for the hollow MOF crystal. The addition of the ditopic linker in MeOH into single crystals of **MOP-macro** produced corresponding hollow single crystals of a $[Cu_{24}(hip)_{24}L_6(H_2O)_{12}]$ formula unit (where L = dabco (**1-macro-h**), pz (**2-macro-h**), and bipy (**3-macro-h**)) (Fig. S6 and S7, ESI†).

Optical photomicrographs of the MOF crystals produced from **MOP-macro** crystal both as a reactant and as a crystalline template indicate that all the MOF crystals are hollow (Fig. 1). When a **1-macro-h** crystal produced from a **MOP-macro** crystal in a MeOH solution of dabco was fragmented, the inner surface of the hollow crystal was exposed in the fragments (Fig. 1a and b).

When hollow crystals of **2-macro-h** and **3-macro-h** in MeOH solutions of pz and bipy, respectively, were fragmented, the release of small crystalline/fine particles encapsulated in the cavity of the hollow crystals was observed (Fig. 1d and f). The encapsulation of the crystalline/fine particles in the inner cavity of the hollow crystal indicates that the size of the particles is larger than the portal dimensions of the corresponding shell MOFs. The particles in the inner cavity are not **MOP-macro** crystals, which are soluble in MeOH, but are probably MOF crystals, which are not soluble in MeOH.

The crystal produced in the process of the transformation from a single crystal of **MOP-macro** to a hollowed single crystal of **1-macro-h** shows that the MOP-to-MOF transformation

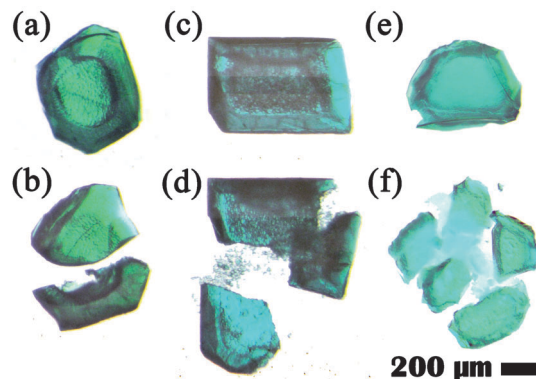


Fig. 1 Optical photomicrographs of **1-macro-h**, **2-macro-h**, **3-macro-h** crystals. (a) A **1-macro-h** crystal and (b) the **1-macro-h** crystal fragmented in two with one piece showing the inner cavity in side view. (c) A **2-macro-h** crystal and (d) the **2-macro-h** crystal fragmented in three in MeOH. (e) A **3-macro-h** crystal and (f) the **3-macro-h** crystal fragmented into several pieces in MeOH.

occurs from the surface to the core of the crystal. The transformation intermediate of a single crystal of **MOP-macro** changes color from cyan to green and from the surface of the crystal to the core while maintaining the core of the crystal as cyan (Fig. 2). In addition, a reactive interface is generated between the shell of the crystalline MOF and the core of the crystalline MOP. While the crystalline MOP core in the fragmented intermediate is immediately dissolved in MeOH (Fig. 2c), the components at the shell interface become small crystalline particles in DMF (Fig. 2d).

A series of isorecticular MOFs of micrometer dimensions, $[Cu_{24}(hip)_{24}L_6(H_2O)_{12}]$ (where L = dabco (**1-micro**), pz (**2-micro**), and bipy (**3-micro**)), with solvent cavities of nanometer dimensions could be prepared *via* stepwise reactions in MeOH using **MOP-macro** as a cuboctahedral MOP precursor (Fig. 3 and Fig. S8, ESI†). The addition of a dabco linker to a MeOH solution of cuboctahedral MOP precursor immediately produced a crystalline powder of **1-micro**. The PXRD pattern of **1-micro** is very similar to that of the reported $[Zn_{24}(mip)_{24}(dabco)_6(H_2O)_{12}]$ MOF (where mip = 5-methylisophthalate) of **ubt** topology (Fig. 3).²² Similar reactions using pz and bipy, respectively, as linkers between the MOP building blocks also produced corresponding isorecticular MOFs of the same **ubt** topology. The PXRD pattern of **2-micro** with pz as a linker is very similar to that of the reported MOF, $[Zn_{24}(mip)_{24}(dabco)_6(H_2O)_{12}]$, because the linker dimension of pz in **2-micro** is almost the same as

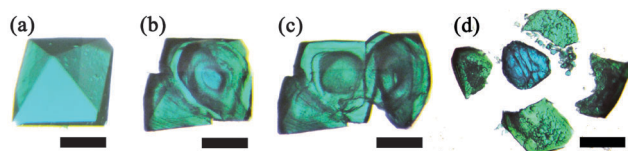


Fig. 2 The optical photomicrographs of a **MOP-macro** crystal and its intermediate in a transformation process to the **1-macro-h** crystal ((a)–(c) scale bar 200 μm, (d) scale bar 500 μm). (a) A **MOP-macro** crystal in DMF, (b) its intermediate, the **MOP-macro** crystal soaked for 20 s in a MeOH solution of dabco, (c) fragments of the intermediate crystal in MeOH, and (d) fragments of another intermediate crystal in DMF.



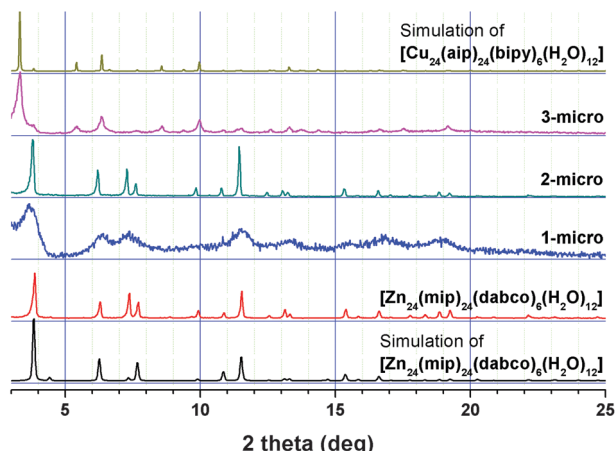


Fig. 3 Comparison of the PXRD patterns of **1-micro**, **2-micro** and **3-micro** with those of $[\text{Zn}_{24}(\text{mip})_{24}(\text{dabco})_6(\text{H}_2\text{O})_{12}]^{22}$ and $[\text{Cu}_{24}(\text{aip})_{24}(\text{bipy})_6(\text{H}_2\text{O})_{12}]^{23}$.

that of dabco in $[\text{Zn}_{24}(\text{mip})_{24}(\text{dabco})_6(\text{H}_2\text{O})_{12}]$ (Fig. 3), and the same MOPs in the two MOFs are interconnected *via* similar lengths of linkers in the same fashion. The strongest reflection peak at about 3.8° in the PXRD pattern of $[\text{Zn}_{24}(\text{mip})_{24}(\text{dabco})_6(\text{H}_2\text{O})_{12}]$ is shifted to about 3.3° for **3-micro**, which indicates an increase of the unit cell dimension with the longer bipy linker between the MOP building blocks (Fig. 3). The PXRD pattern of **3-micro** matches well with that of the reported $[\text{Cu}_{24}(\text{aip})_{24}(\text{bipy})_6(\text{H}_2\text{O})_{12}]$ (where aip = 5-aminoisophthalate) of the same **ubt** topology with the same bipy linker.²³

Hollow single crystalline MOFs of micrometer dimensions could be prepared by reacting **MOP-micro** crystals of micrometer dimensions with ditopic linkers in MeOH (Fig. S9, ESI†). Quick addition of a MeOH solution of dabco to **MOP-micro** crystals wet with a trace amount of DMF produced micro-sized hollow MOF crystals, **1-micro-h** (Fig. S10, ESI†). The shell of a hollow MOF is not an aggregate of small non-hollow single crystals, but is a single crystalline MOF. SEM images of the hollow **1-micro-h** crystals show that most have the appearance of ordinary non-hollow crystals (Fig. S10c and d, ESI†) while some have holes at their surface (Fig. S11b, ESI†). By contrast, more fragmented hollow MOFs are observed in hollow **2-micro-h** crystals (Fig. S10e and S11c, ESI†). The inner surface of the fragmented hollow MOF is rougher than the outer surface. While the outer surface is part of the single crystalline MOF, the inner surface appears to be an aggregate of the smaller crystalline particles. Most hollow **3-micro-h** crystals also have the appearance of ordinary non-hollow crystals (Fig. S10g and h, ESI†), while a few have holes at their surface (Fig. S11d, ESI†). The crystals have octahedral morphology and have smooth and clean outer surfaces. The **3-micro-h** crystals are quite rigid. Even after sonication, most **3-micro-h** crystals maintain their morphology and have clean outer surfaces (Fig. S12, ESI†). The shell of the hollow **3-micro-h** is not an aggregate of smaller crystalline particles, but is a single crystalline particle. Sonication of hollow **3-micro-h** single crystals produced more fragmented hollow crystals and showed some interesting characteristics of the hollow MOFs, such as roughness of their inner surface and the approximate thickness of the fragmented shell MOF crystals (Fig. S12c, ESI†).

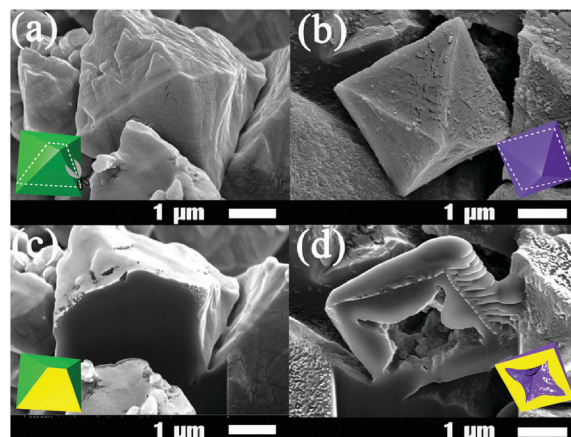


Fig. 4 SEM images of (a) a nonhollow **MOP-micro** crystal and (b) a hollow **3-micro-h** MOF crystal obtained from nonhollow **MOP-micro** crystal. FIB-SEM images of (c) the **MOP-micro** crystal sliced by using FIB milling and (d) the **3-micro-h** crystal sliced by using FIB milling after carbon coating.

The hollow nature of the micro-h crystals was confirmed by using focused ion beam-scanning electron microscopy (FIB-SEM) (Fig. 4 and Fig. S13, ESI†). To see the inner cavity of a hollow MOF crystal, the crystal was sliced using Ga ion milling. In contrast to the FIB-SEM image of the sliced non-hollow **MOP-micro** crystal (Fig. 4c), that of the sliced hollow **3-micro-h** MOF crystal (Fig. 4d) clearly shows its inner cavity and confirms the hollow nature of the seemingly non-hollow-like MOF crystal.

The total volume of the **3-micro-h** product is much larger than that of the starting **MOP-micro** reactant (Fig. 5). The volume increase in the **3-micro-h** crystals is not only the result of expansion of the unit cell volume of the **3-micro** crystal compared with that of the **MOP-micro** crystals, but is also because of the formation of a hollowed cavity of micrometer dimensions (Fig. 5).

Unlike other approaches for hollow MOFs, single crystalline hollow MOFs could be obtained by using a MOP single crystal as a sacrificial template. The MOP crystal serves not only as a hard template, but also as a reactant for single crystalline hollow MOFs (Scheme 2). At the initial stage, the MOP building units weakly interacting with each other at the surface of a MOP crystal diffuse outward into the methanol solution containing ditopic linkers and generate a reactive interface containing both the MOP building units and the ditopic linkers. The reaction at the

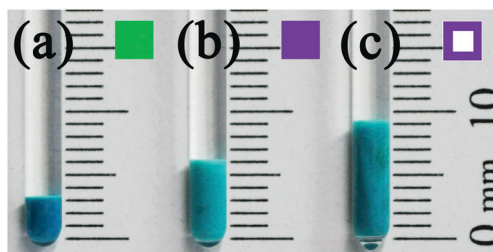
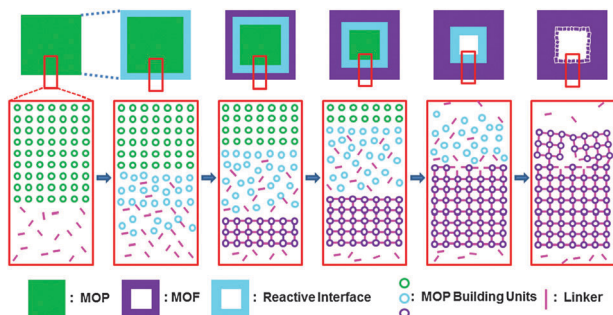


Fig. 5 (a) The volume of **MOP-micro** crystals, (b) the volume of the **3-micro** crystals obtained from the **MOP-micro** crystals, and (c) the corresponding volume of the **3-micro-h** crystals obtained from the same amount of the **MOP-micro** crystals.





Scheme 2 The proposed formation mechanism for a single crystalline hollow MOF using a MOP single crystal as a sacrificial hard template.

reactive interface generates a thin crystalline microporous MOF shell, which accompanies the expansion of the overall dimension of the crystal and the subsequent formation of a new reactive interface. Because the small ditopic linkers in the MeOH solvent can still interpenetrate the microporous shell of the crystalline MOF, a new reactive interface with loosely bound MOP building units and the ditopic linkers can be generated between the crystalline MOF shell and the crystalline MOP core.

The microporosity of the crystalline MOF shell results in a difference in the diffusion rates of the ditopic linkers and the MOP building units. While the ditopic linkers are sufficiently small enough to diffuse into the crystal through the microporous crystalline MOF shell, the MOP building units are too large to diffuse out of the crystal through the microporous crystalline MOF shell. Consequently, the growth of the crystalline MOF shell only proceeds inward with the progressive movement of the reactive interface from the surface to the core of the crystal. As the formation of the crystalline MOF shell proceeds, the thickness of the MOF shell increases and the size of the crystalline MOP decreases. When all the MOP units at the core of the crystal are expended, single crystalline hollow MOFs with hollow cavity dimensions of several micrometers and several hundred micrometers are generated depending on the size of the MOP single crystals. The formation of the hollow structure and its dimensions are dependent on several factors such as the size of the MOP single crystal, the kind of ditopic linker and its concentration, and the solvent employed. Those factors also affect the formation of small microcrystals with varying dimensions at the microscopic/macroscale of the single crystalline hollow MOF.

In summary, single crystalline hollow MOF structures were obtained by using a MOP single crystal as a sacrificial hard template. The formation of the microporous MOF shell in the hollow MOF occurs *via* the diffusion of the cuboctahedral Cu-MOP building units at the surface of the templating MOP

single crystal outward to the MeOH solution containing the linker. While cuboctahedral Cu-MOP building units larger than the pore dimensions of the microporous MOF shell could not diffuse outward through the micropores of the MOF shell from the sacrificial MOP single crystal, linkers smaller than the pore dimension can diffuse inward through the micropores of the MOF shell to the cuboctahedral Cu-MOP building unit, which leads to the formation of the crystalline hollow MOFs. The dimensions of the hollow MOFs can be controlled by the dimensions of the sacrificial MOP single crystals. Not only hollow MOFs of the order of hundred micrometer dimensions, but also those of the order of micrometer dimensions could be obtained.

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