



# Magnesium Nanocrystals Embedded in a Metal–Organic Framework: Hybrid Hydrogen Storage with Synergistic Effect on Physi- and Chemisorption\*\*

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Hydrogen is considered to be a promising energy carrier for the future. In order to use it as a fuel, a compact, safe, and efficient hydrogen storage system should be developed. Porous metal–organic frameworks (MOFs) have attracted great attention as potential hydrogen storage materials,<sup>[1]</sup> because some MOFs can store large amount of H<sub>2</sub> (> 7 wt %) at 77 K and high pressures.<sup>[2]</sup> However, at room temperature, the H<sub>2</sub> storage capacities of MOFs drop to less than 1 wt % because interaction energies between the frameworks and H<sub>2</sub> are very low (4–8 kJ mol<sup>−1</sup>). There have been efforts to enhance the H<sub>2</sub> storage capacities of MOFs at ambient temperature by methods such as tuning the ligand,<sup>[3]</sup> generation of open metal sites,<sup>[4]</sup> and embedding palladium nanoparticles.<sup>[5]</sup> Despite these modifications, the improvements were not very satisfactory, and none of the materials have yet met the target of the US Department of Energy (DOE) of 2017.

In search for a highly efficient material for H<sub>2</sub> storage, we have investigated the possibilities of hybrid materials that would be capable of storing H<sub>2</sub> by both physical adsorption and chemisorption. In particular, we have been interested in composite materials that consist of a MOF and magnesium nanoparticles. Magnesium chemisorbs H<sub>2</sub> at 773 K and 200 atm in the presence of MgI<sub>2</sub> catalyst.<sup>[6]</sup> Magnesium powder of 50–100 μm size made by ball milling slowly absorbs H<sub>2</sub> at 673 K and 10 bar.<sup>[7]</sup> The resulting magnesium hydride has a quite high dehydrogenation enthalpy (≈ 75 kJ per mol H<sub>2</sub>),<sup>[8]</sup> and therefore heavy-metal catalysts should be used to

reduce the H<sub>2</sub> desorption temperature and improve kinetics, but they still do not allow effective dehydrogenation.<sup>[9]</sup> In the presence of heavy-metal catalysts, the H<sub>2</sub> absorption and release temperatures decrease as a result of destabilization of the metal hydride when the particle size of magnesium is reduced.<sup>[10]</sup>

Nanosized magnesium particles have been synthesized by a variety of methods, such as ball-milling,<sup>[11]</sup> condensation of magnesium metal vapor,<sup>[12]</sup> plasma metal reaction,<sup>[13]</sup> infiltration of melted Mg in carbon material,<sup>[14]</sup> and sonoelectrochemistry<sup>[15]</sup> or chemical reduction of Mg precursors.<sup>[16]</sup> Some of the disadvantages of these methods are that they require a long process time, extremely high temperatures (898–1203 K), and electrically or chemically harsh conditions, and yet they provide inhomogeneous size distributions.

Herein we report a simple method for fabrication of hexagonal-disk-shaped magnesium nanocrystals (Mg NCs) within a MOF. The fabrication of Mg nanocrystals inside MOFs is unprecedented. Previously, small Ag, Au, Ni, Pd, Ru, or Pt nanoparticles (size, 1.4–10 nm) were prepared in MOFs by immersing redox-active MOFs in the metal-ion solutions<sup>[5,17]</sup> or by reducing the metal precursors deposited in the MOFs.<sup>[18]</sup> The present nanocomposite is made by the thermal decomposition of air-sensitive bis-cyclopentadienyl magnesium (MgCp<sub>2</sub>) vapor in a MOF, leading to the deposition of Mg NCs in the MOF, a method that is also unprecedented for production of nanosized Mg. The resulting Mg NCs@MOF is a hybrid hydrogen storage material that stores H<sub>2</sub> by both physical adsorption and chemisorption, exhibiting a synergistic effect to increase the isosteric heat of H<sub>2</sub> physisorption and decrease the temperatures for chemisorption/desorption of H<sub>2</sub>.

Yellow crystals of [Zn<sub>4</sub>O(atb)<sub>2</sub>]·22DMF·9H<sub>2</sub>O (SNU-90), in which atb is aniline-2,4,6-tribenzoate, were synthesized by heating a mixture of Zn(NO<sub>3</sub>)<sub>2</sub> and H<sub>3</sub>(atb) in dimethylformamide (DMF). The X-ray single-crystal structure of SNU-90 indicated a non-interpenetrated (6,3)-connected net of a *qom* topology, similar to MOF-177<sup>[19]</sup> (see the Supporting Information, Figures S1 and S2). When the guest solvent molecules in SNU-90 were removed by treatment with supercritical CO<sub>2</sub>,<sup>[2b,20]</sup> a guest-free [Zn<sub>4</sub>O(atb)<sub>2</sub>] (SNU-90', NH<sub>2</sub>-MOF-177) was obtained.

Vapor of bis(cyclopentadienyl) magnesium (MgCp<sub>2</sub>) was deposited in SNU-90' at 80 °C, and the resulting MgCp<sub>2</sub>@SNU-90' was thermally decomposed at 200 °C under an argon atmosphere followed by evacuation, which gave rise to Mg NCs embedded in SNU-90'. Thus far it was unknown that the thermal decomposition of MgCp<sub>2</sub> results in

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**Table 1:** N<sub>2</sub> and H<sub>2</sub> gas uptake data in SNU-90' and various samples of Mg@SNU-90'.

| Compound    | mol <sub>Mg</sub> /mol <sub>Zn</sub> (wt %) | N <sub>2</sub> uptake<br>[cm <sup>3</sup> g <sup>-1</sup> ] | Surface area<br>[m <sup>2</sup> g <sup>-1</sup> ] |                     | Pore volume<br>[cm <sup>3</sup> g <sup>-1</sup> ] | H <sub>2</sub> uptake [wt %] |                                                                      | Q <sub>st</sub><br>[kJ mol <sup>-1</sup> ] |
|-------------|---------------------------------------------|-------------------------------------------------------------|---------------------------------------------------|---------------------|---------------------------------------------------|------------------------------|----------------------------------------------------------------------|--------------------------------------------|
|             |                                             |                                                             |                                                   |                     |                                                   | at 1 atm                     | at high pressure                                                     |                                            |
| SNU-90'     | n.a.                                        | 1135                                                        | 4244 <sup>[a]</sup>                               | 4914 <sup>[b]</sup> | 1.64 <sup>[c]</sup>                               | 1.21 (77 K)<br>0.74 (87 K)   | 8.81 (77 K, 75 bar)<br>0.45 (298 K, 80 bar)                          | 4.55                                       |
| Mg@SNU-90'a | 0.15 (1.26)                                 | 1104                                                        | 4154 <sup>[a]</sup>                               | 4757 <sup>[b]</sup> | 1.47 <sup>[c]</sup>                               | 1.24 (77 K)<br>0.65 (87 K)   | 8.74 (77 K, 89 bar)<br>0.54 (298 K, 90 bar)                          | 5.68                                       |
| Mg@SNU-90'b | 0.85 (6.52)                                 | 559                                                         | 2056 <sup>[a]</sup>                               | 2373 <sup>[b]</sup> | 0.84 <sup>[c]</sup>                               | 0.72 (77 K)<br>0.40 (87 K)   | 0.29 (473 K, 30 bar)                                                 | 7.24                                       |
| Mg@SNU-90'c | 1.40 (10.5)                                 | 378                                                         | 1371 <sup>[a]</sup>                               | 1581 <sup>[b]</sup> | 0.36 <sup>[c]</sup>                               | 0.60 (77 K)<br>0.47 (87 K)   | 0.20 (323 K, 80 bar)<br>0.24 (415 K, 40 bar)<br>0.71 (473 K, 30 bar) | 11.6                                       |

[a] BET surface area. [b] Langmuir surface area, and [c] Pore volume estimated by using the Dubinin–Radushkevich (DR) equation. n.a. = not available.

nanosized Mg, although it was reported that reduction of MgCp<sub>2</sub> with Li<sup>+</sup>(naphthalenide<sup>-</sup>) in a solution of polymethylmethacrylate in tetrahydrofuran afforded round-shaped Mg nanoparticles.<sup>[16]</sup> As for the mechanism of the thermal decomposition, formation of metal and two C<sub>5</sub>H<sub>5</sub> radicals that might further dissociate or associate to the complex products, was predicted as an initial step based on the theoretical calculation.<sup>[21]</sup> However, in order to verify the mechanism in our case, further research should be carried out. By changing the reaction conditions during the deposition process of MgCp<sub>2</sub>, such as the duration time and whether or not a prevacuum state was applied to the reaction vessel, we could control the loaded amounts of MgCp<sub>2</sub> in the MOF and obtained several samples embedded with different amounts of magnesium nanocrystals (Supporting Information, Table S1).

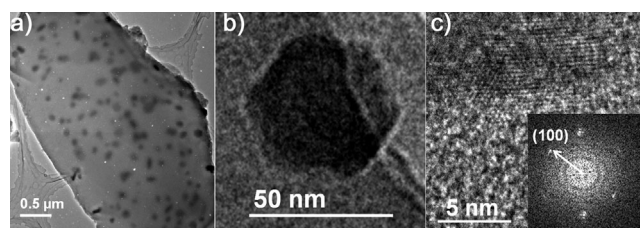
The samples, Mg@SNU-90'a, Mg@SNU-90'b, and Mg@SNU-90'c, were loaded with 1.26 wt %, 6.52 wt %, and 10.5 wt % of Mg NCs, respectively, as determined by inductively coupled plasma atomic emission spectroscopy (ICP-AES) (Table 1). The NMR spectra indicated the absence of cyclopentadiene (Cp) in the resulting nanocomposites (Supporting Information, Figure S3).

The HRTEM images for various samples of Mg@SNU-90' showed hexagonal-disk-shaped magnesium nanocrystals,<sup>[22]</sup> which have a diagonal length of hexagon ranging 44–88 nm (average (60 ± 18) nm) with a thickness ranging 16–61 nm (average (37 ± 12) nm; see Figure 1 and Figure S4 in the Supporting Information). Although it was reported that hexagonal-disk-shaped Mg NCs could be formed when Mg powder was heated at temperatures higher than 500 °C,<sup>[22b]</sup> in the present work, they are formed by heating Mg(Cp)<sub>2</sub>@MOF

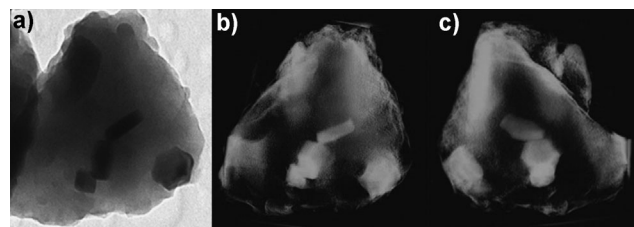
at only 200 °C. The formation of Mg NCs that are much bigger than the channel size (12.9 Å × 9.6 Å) of the MOF is attributed to the agglomeration of nanoparticles by heat during the thermal decomposition. Alternatively, it is simply caused by the degradation of the MOF under electron-beam irradiation during the observation by HRTEM.<sup>[23]</sup> The selected-area electron diffraction (SAED) patterns of Mg NCs embedded in SNU-90' showed lattice fringes separated by 2.7871 Å, which is in agreement with the (100) *d*-spacing of metallic Mg (2.7782 Å, JCPDS 04-0770). The size distribution of the Mg NCs was found to be very similar in all of the samples, even though there were significantly different amounts of Mg NCs embedded in the MOF. When Mg@SNU-90' was exposed to air for three weeks, the shape of the Mg NCs changed to a star shape that has a lattice fringe with a separation of 2.1101 Å, which corresponds to the (200) *d*-spacing of MgO (2.1061 Å, JCPDS 89-7746; Supporting Information, Figure S4). XPS data of Mg@SNU-90' indicated that Mg<sup>0</sup> and Zn<sup>II</sup> coexisted in the solid (Supporting Information, Figure S5).

To verify that the Mg NCs were embedded inside the channels of the MOF, we performed electron tomography. From the TEM images that were collected with a 1° interval with the tilting angle of +40°–40°, 3D images were constructed by using the IMOD program.<sup>[24]</sup> When the sample was tilted, the shape of a Mg NC changed from hexagon to rectangle and from rectangle to hexagon, thus indicating that the Mg NCs were indeed embedded in the MOF (Figure 2 and Supporting Video).

The powder X-ray diffraction (PXRD) patterns showed that the structure of SNU-90' was maintained even after the formation of Mg NCs, the sizes of which were much larger than the channel size (12.9 Å × 9.6 Å) of the MOF (Supporting Information, Figure S6). To understand this observation,



**Figure 1.** High-resolution transmission electron microscopy (HRTEM) images. a) Mg@SNU-90'b. b) A hexagonal-shaped Mg nanocrystal and c) its edge image showing lattice fringes with a separation of 2.7871 Å, which corresponds to (100) *d*-spacing of Mg<sup>0</sup> (2.7782 Å, JCPDS 04-0770). Inset: selected-area electron diffraction (SAED) pattern.



**Figure 2.** TEM and electron tomography images of Mg@SNU-90'b. a) TEM image. b) and c) Tomographically reconstructed images of Mg NCs embedded in SNU-90' at different angles.

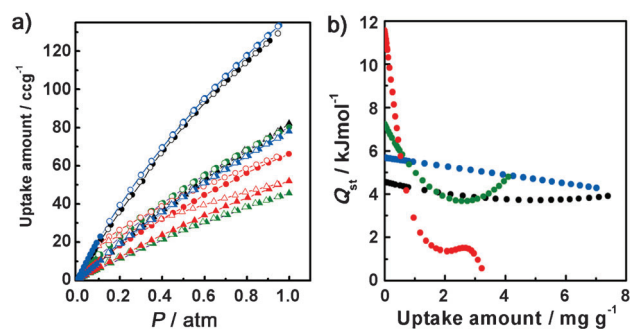
we calculated the ratio of the volumes of formed Mg NCs and the network skeleton for Mg@SNU-90'c. Assuming that all Mg NCs destroyed the network skeleton, the Mg NCs would destroy a maximum 4.6 % of the network skeleton by volume. This value is too small to alter the PXRD pattern of the MOF. It was also previously reported that metal nanoparticles much bigger than the cavity sizes of the hosts were impregnated by maintaining the network structures.<sup>[5,17]</sup> The PXRD peaks of crystalline Mg should generally appear at  $2\theta = 32.2$ ,  $34.4$ , and  $36.6^\circ$ , but we could observe only a peak at  $36.6^\circ$ , which is the strongest peak of Mg (Figure S7). On exposure to air for two days, however, the PXRD pattern showed the peaks corresponding to MgO, with retention of the framework structure of SNU-90'.

The adsorption–desorption isotherms were measured for N<sub>2</sub> gas at 77 K and for H<sub>2</sub> gas at various temperatures on SNU-90', Mg@SNU-90'a, Mg@SNU-90'b, and Mg@SNU-90'c (Figure 3 Table 1). The N<sub>2</sub> gas sorption isotherms of the four samples were of type I, which is characteristic for the microporous materials (Supporting Information, Figure S8.). Despite the presence of the NH<sub>2</sub> group in the ligand of SNU-90', the surface area, pore volume, and H<sub>2</sub> adsorption capacities of SNU-90' were similar to those of MOF-177.<sup>[19]</sup> Gas sorption data for various samples of Mg@SNU-90' showed that as the amount of Mg increased, the BET surface area, pore volume, and H<sub>2</sub> uptake capacity at 77 K and 1 atm decreased, because Mg NCs occupied the surface and space of the pores in the MOF. At 298 K and high pressure, however, the H<sub>2</sub> uptake in Mg@SNU-90'a increased by 20 %, to 0.54 wt % from 0.45 wt % in pristine SNU-90', thus suggesting that Mg NCs provide a positive effect on H<sub>2</sub> adsorption at 298 K (Supporting Information, Figure S9). The zero-coverage isosteric heats of the H<sub>2</sub> adsorption, which were estimated from the H<sub>2</sub> adsorption isotherms measured at 77 K and 87 K, increased as the amount of Mg increased, up to 11.6 kJ mol<sup>-1</sup> for Mg@SNU-90'c from 4.55 kJ mol<sup>-1</sup> for SNU-90' (Table 1, Figure 3).

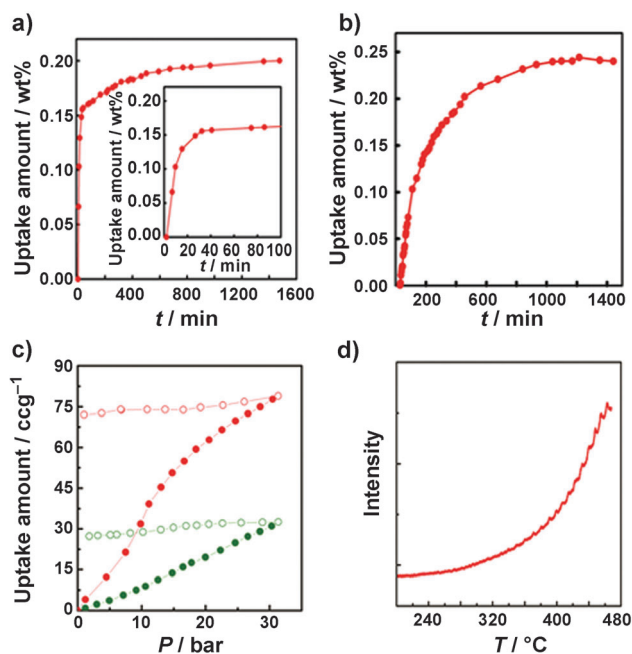
In order to observe the chemisorption ability of the embedded Mg NCs, we also measured the H<sub>2</sub> uptake in Mg@SNU-90'c at 323 K, 415 K, and 473 K under high pressures (Figure 4). Contrary to the physisorbed MOFs, in

which H<sub>2</sub> uptake decreases at the elevated temperature, the H<sub>2</sub> chemisorption capacities of the present material increase as the temperature is raised. The H<sub>2</sub> absorption capacities of Mg@SNU-90'c were 0.20 wt % at 323 K under the H<sub>2</sub> pressure of 80 bar, 0.24 wt % at 415 K under the H<sub>2</sub> pressure of 40 bar. At 473 K under 30 bar, the H<sub>2</sub> uptake capacity of Mg@SNU-90'c became 0.71 wt % (volumetric H<sub>2</sub> storage capacity, 3.1 g L<sup>-1</sup>). This value is remarkably higher than that of Mg@SNU-90'b under the same conditions (Table 1), despite the former exhibiting a much lower surface area and pore volume because of the heavily loaded Mg. The fact that the H<sub>2</sub> uptake enhances with elevated temperatures and the increased amount of Mg NCs confirms that the H<sub>2</sub> uptakes in Mg@SNU-90' at 323 K, 415 K, and 473 K are results of chemisorption. These chemisorption temperatures are significantly lower (by > 200 K) than that (673 K under 10 bar) of bare Mg powder of 50–100  $\mu$ m in size.<sup>[7]</sup> If the H<sub>2</sub> uptake capacity of Mg alone is estimated from the data, it is 7.5 wt % at 473 K and 30 bar. Considering that the H<sub>2</sub> chemisorption capacity of pure Mg is 7.66 wt %, 99 % of the Mg NCs in the sample chemisorb H<sub>2</sub>, which is much better than any form of Mg previously reported, such as Mg nanoparticles incorporated in a polymer<sup>[16]</sup> and bare Mg powder.<sup>[25]</sup>

The H<sub>2</sub> desorption properties of Mg@SNU-90'c were verified by the temperature-programmed desorption mass spectroscopy (TPD-MS) analysis (Figure 4d). The TPD-MS results indicated that H<sub>2</sub> was desorbed at  $T > 523$  K and 1 atm. It should be noted that the signal intensities of hydrogen atoms were exactly twice as strong as those of H<sub>2</sub> molecules (Supporting Information, Figure S10). This observation sug-



**Figure 3.** Physical adsorptions of H<sub>2</sub> in SNU-90' (black), Mg@SNU-90'a (blue), Mg@SNU-90'b (green), and Mg@SNU-90'c (red). a) The H<sub>2</sub> adsorption isotherms at 77 K (circles) and 87 K (triangles). Filled shape: adsorption, open shape: desorption. b) Isosteric heats of the H<sub>2</sub> adsorption. The samples of Mg@SNU-90'a, Mg@SNU-90'b, and Mg@SNU-90'c contain 1.26 wt %, 6.52 wt %, and 10.5 wt %, respectively, of magnesium nanocrystals.



**Figure 4.** Chemical absorption of H<sub>2</sub> in Mg@SNU-90'c. a) H<sub>2</sub> absorption kinetics at 325 K and 80 bar. Inset: data with a magnified time scale. b) H<sub>2</sub> absorption kinetics at 415 K and 40 bar. c) H<sub>2</sub> absorption isotherms at 473 K in Mg@SNU-90'b (green) and Mg@SNU-90'c (red). Filled shapes: adsorption; open shapes: desorption. d) Temperature-programmed desorption mass spectroscopy (TPD-MS) data;  $m/e = 2$ , measured under argon.



gests that all desorbed H<sub>2</sub> sources were MgH<sub>2</sub>, not from the MOF at all. The TG mass loss of Mg@SNU-90'c after H<sub>2</sub> absorption is also in good agreement with the TPD-MS data (Supporting Information, Figure S11).

After the H<sub>2</sub> chemisorption at 473 K and 30 bar, the HRTEM images of Mg@SNU-90'c indicated that the crystal morphology of Mg was maintained but the crystal size was remarkably increased. The SAED pattern indicated the lattice fringe with a separation of 2.247 Å, which is in good agreement with the (110) *d*-spacing of β-MgH<sub>2</sub> (2.257 Å, JCPDS 35-1185, Supporting Information, Figure S12). The MgH<sub>2</sub>@SNU-90' shows the crystal size (diagonal length of hexagon) of average (142 ± 41) nm and thickness of average (83 ± 18) nm (Supporting Information, Figure S12), circa 2.3 times as large as those of Mg NCs. Considering that the volume of Mg should be increased by 30% on hydride formation,<sup>[26]</sup> this excessive expansion during the H<sub>2</sub> chemisorption processes at high temperature and pressure can be explained by the three-dimensional Ostwald ripening, in which the larger crystals take up the mobile atoms dissociated from the smaller crystals.<sup>[27]</sup> In addition, degradation of the MOF by electron beam during the TEM measurement plays a role in increasing the size of the MgH<sub>2</sub> crystal. Even after the formation of MgH<sub>2</sub>, the organic ligands of SNU-90' were not hydrogenated, as evidenced by the <sup>1</sup>H NMR spectra measured for the resulting nanocomposite that was dissolved in the mixture of DCl and [D<sub>6</sub>]Me<sub>2</sub>SO (Supporting Information, Figure S3). When MgH<sub>2</sub>@SNU-90' was exposed to air, the PXRD pattern data showed new peaks at 2θ = 36.8 and 42.8 deg, which correspond to MgO, and a new peak at 2θ = 38.0 deg that corresponds to Mg(OH)<sub>2</sub>, (Supporting Information, Figure S6).

In conclusion, we have prepared for the first time Mg nanocrystals in a MOF, and showed that Mg NCs@MOF is a hybrid-hydrogen-storage material that has both physical adsorption and chemisorption properties, and exhibits synergistic effects to increase isosteric heat of the H<sub>2</sub> physisorption and to decrease the temperatures for chemisorption/desorption of H<sub>2</sub>. The present hybrid-hydrogen-storage material suggests a new strategy for H<sub>2</sub> storage and transport for the future. To develop the materials that are able to reach the DOE target, we are trying to further reduce the size of the Mg NCs and adjust the amount of loaded Mg in the selected physisorbed porous materials, such as light-weighted MOFs or organic COFs.<sup>[28]</sup> In addition, we are trying to develop new methods to protect the material from oxidation through air.

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