Hydrazide-Linked Covalent Organic Frameworks for Water Harvesting

Ha L. Nguyen,†‡,‡,‡,‡ Cornelius Gropp,†‡,‡,‡ Nikita Hanikel,† Anna Möckel,† Alicia Lund,‡ and Omar M. Yaghi†‡,‡,

†Department of Chemistry, University of California–Berkeley; Kavli Energy Nanoscience Institute at UC Berkeley; and Berkeley Global Science Institute, Berkeley, California 94720, United States
‡Joint UAEU–UC Berkeley Laboratories for Materials Innovations, United Arab Emirates University, Al-Ain 15551, United Arab Emirates
‡Department of Chemistry, University of California–Berkeley, Berkeley, California 94720, United States

KEYWORDS: covalent organic frameworks, postsynthetic modification, oxidation, cyclization, water harvesting.

ABSTRACT: We report a postsynthetic strategy and its implementation to make covalent organic frameworks (COFs) with irreversible hydrazide linkages. This involved the synthesis of three 2D and 3D hydrazine-linked frameworks and their oxidation. The linkage synthesis and functional group transformation—hydrazine and hydrazide—was evidenced by $^{15}$N multi-CP-MAS NMR. In addition, the isothermal water uptake profiles of these frameworks were studied, leading to the discovery of one hydrazide-linked COF suitable for water harvesting from air in arid conditions. This COF displayed characteristic S-shaped water sorption profiles, a steep pore-filling step below 18% relative humidity at 25 °C, and a total uptake capacity of 0.45 g g$^{-1}$. We found that even small changes made on the molecular level can lead to major differences in the water isotherm profiles, therefore pointing to the utility of water sorption analysis as a complementary analytical tool to study linkage transformations.

INTRODUCTION

Multi-step organic synthetic methods have been successfully translated to the solid state. This practice has become especially attractive for the synthesis of new covalent organic frameworks (COFs). Their hallmark features, porosity and crystallinity, allow chemical transformations to be done on the framework and enable rigorous analysis of concurrent structural changes through diffraction and spectroscopic techniques.

The synthesis of COFs relies on an initial reversible step to guide the covalent assembly of organic building units into extended and crystalline structures. This step can be followed by one or more postsynthetic transformations, leading to structural complexity and function of the framework. This has been demonstrated for a number of imine-linked COFs, but hydrazide moieties have not been reported yet.

Here, we report three 2D and 3D hydrazide-linked COFs following postsynthetic oxidation of the hydrazine-linking units (Scheme 1). Considering the rather hydrophilic nature of the hydrazine and hydrazide linkages, we chose to study the effect of the synthetic transformations with water vapor sorption analysis. This not only allowed us to explore new sorbents for water harvesting from air, but also provided us with additional insight into the micro- and macroscopic changes occurring on the framework.

Scheme 1. Synthetic strategy for hydrazide-linked COFs. This strategy was applied to a molecular model system and two 2D (hcb, sql) and one 3D (dia) COFs.

Our synthetic strategy towards hydrazide-linked COFs is based on the reversible formation of the hydrazine-linkage, followed by postsynthetic oxidation to the hydrazide. (Scheme 1). We first demonstrated the feasibility of this route on a molecular model system. Then, we explored its transferability to the solid-state by synthesizing two 2D COFs with honeycomb (hcb) and square-lattice (sql) topologies and one 3D COF with diamond (dia) topology. In total, six COFs were synthesized, four of which were hitherto unreported, and comprised of hydrazine and hydrazide linkages—the latter presents a linkage that has not been reported in COF chemistry yet. The COFs were analyzed by $^{15}$N multiple cross-polarization magic-angle spinning spectroscopy.
Scheme 2. Overview of the synthesis of (a) 2D hcb, (b) 2D sql, and (c) 3D dia hydrazide-linked COFs. The synthesis of hydrazine-linked COFs is detailed in the SI.

To demonstrate the transferability from solution to the solid state, we first synthesized two 2D and one 3D hydrazine-linked COFs (Scheme 2). AB-COF\textsuperscript{20,21} was produced from the trigonal-planar 1,3,5-benzenetrialdehyde (BTA) and linear hydrazine (Hy), and crystallized in the hcb topology with the reticular formula \([\text{BTA},_3(\text{Hy}),_3]\)\textsubscript{hydrazine}. Py-COF-2D\textsuperscript{22} is comprised of the square-planar 1,3,6,8-tetrakis(p-formylphenyl)pyrene (TFPPy) unit and Hy and crystallized in the sql topology with the reticular formula \([(\text{TFPPy})(\text{Hy}),_2]\)\textsuperscript{hydrazine}. Lastly, COF-670-hydrazine was synthesized from the tetrahedral 4,4',4'',4'''-methanetetrayltetra benzaldehyde (MTBA) and Hy and crystallized in the dia topology with the reticular formula \([(\text{MTBA})(\text{Hy}),_2]\)\textsuperscript{hydrazine} (Scheme 2). While AB-COF and Py-COF-2D have been previously reported,\textsuperscript{20,21} a hydrazine-linked 3D COF, COF-670-hydrazine, has not been reported to date.

We first explored the oxidation of the hydrazine-linked AB-COF by subjecting it to our reported oxidation (multi-CP-MAS NMR),\textsuperscript{19} giving a quantitative readout of the chemical transformations. Fourier-transform infrared (FT-IR), solid- and solution-state nuclear magnetic resonance (NMR) spectroscopies, elemental analysis (EA), thermogravimetric analysis (TGA), powder X-ray diffraction (PXRD), and nitrogen sorption substantiated the structural and compositional integrity of the frameworks.

RESULTS AND DISCUSSION

To show the viability of our postsynthetic strategy, we first synthesized a hydrazine molecular model compound, 1,2-benzyldenedehydrazone, from benzaldehyde and hydrazine with quantitative yield (Scheme 1).\textsuperscript{19} The hydrazine compound was subsequently oxidized to afford the hydrazide molecular model using previously reported conditions for the conversion of imines to amides in COFs (see Supporting Information, SI, Section S2).\textsuperscript{5}
conditions involving the use of NaClO₃ as oxidant in aqueous acidic medium in the presence of 2-methyl-2-butene over three days to obtain COF-480-hydrazide (see SI, Section S2). FT-IR spectroscopic traces of the product indicated attenuation of the imine band at 1625 cm⁻¹ corresponding to the hydrazine linkages and an emerging band at 1674 cm⁻¹, which was assigned to the carbonyl stretching vibration of the hydrazide moiety (see SI, Section S3). The solid-state ¹⁵N multi-CP-MAS NMR spectroscopy, performed on a 50% ¹⁵N-labeled sample, further corroborated the transformation with a new peak at 171.7 ppm. However, no quantitative conversion was observed, despite prolonged reaction times or increased amount of oxidation reagent (see SI, Section S2). This led us to the conclusion that only partial oxidation was achieved—an observation that we further substantiated with an ¹⁵N NMR experiment on the hydrazide model system (see SI, Section S4). Multi-CP-MAS NMR on the ¹⁵N labeled COF samples allowed us to quantify the conversion, which amounted to 9.2% (see SI, Section S4). Here, the resonances corresponding to the functionalities were fit to Gaussian line shapes and their relative areas were compared. This analysis correlated well with the atomic composition determined by EA (see SI, Section S2).

TGA under nitrogen and air flow revealed an onset in decomposition for the pristine and the oxidized samples at around 380 ºC.

PXRD analysis of the microcrystalline powder confirmed retention of crystallinity after oxidation and did not reveal additional diffraction peaks associated with hydrolyzed starting materials or additives (see SI, Section S6). The PXRD pattern of AB-COF shows high-intensity peaks at 6.8°, 11.8°, and 26.4° 2θ values, associated with the (100), (1-10), and (002) lattice planes. After oxidation, the broad peak indexed as the lattice plane (002) shifts to a higher 2θ value (26.6°), resulting from changes in the interlayer stacking of the structure.

N₂ sorption analysis at 77 K of the pristine AB-COF and COF-480-hydrazide demonstrated permanent porosity and revealed a Type I isotherm, indicating a micropore structure. The Brunauer–Emmett–Teller (BET) surface areas were calculated to be 1209 m² g⁻¹ and 989 m² g⁻¹, respectively. The pore size distribution, estimated from the N₂ isotherms and calculated by density functional theory (DFT) using the cylinder geometry, indicated comparable pore sizes of around 11 Å in diameter (see SI, Section S7).

To gain further insight into the impact of the chemical transformations on the water sorption behavior, we measured the water sorption isotherms at 15 ºC, 25 ºC, and 35 ºC (Figure 1a, b). AB-COF displays an S-shaped water sorption isotherm with a steep pore-filling step at around 23% relative humidity (RH) at 25 ºC (Figure 1a). The maximal uptake at P/Pₕ sat = 0.95 reaches a total uptake capacity of 0.45 g g⁻¹. After oxidation, the water sorption isotherm of COF-480-hydrazide substantially shifts to lower RH with a steep pore-filling step at 8%—thus extending the RH range at which COF-480-hydrazide can operate to more arid conditions. Importantly, the COF preserves its total water uptake capacity of around 0.45 g g⁻¹ at P/Pₕ sat = 0.95 (25 ºC).

Furthermore, the water sorption isotherm profiles are retained at 15 ºC and 35 ºC, indicating robust sorption behavior under varying temperatures. The increased water uptake capacity at lower RH upon oxidation indicated stronger water–framework interactions during the pore-filling step. To evaluate the strength of these interactions, the isosteric heat of water adsorption (Qₐ) was calculated with the Clausius–Clapeyron equation using the water isotherms at 15 ºC, 25 ºC, and 35 ºC. The Qₐ was estimated to be 49 kJ mol⁻¹ for the pristine AB-COF and 50.5 kJ mol⁻¹ for the oxidized COF-480-hydrazide. A ΔQₐ = 1.5 kJ mol⁻¹ upon oxidation likely results from the stronger hydrogen-bonding donor and acceptor sites in the

Figure 1. Water vapor sorption analysis of AB-COF (a) and COF-480-hydrazide (b) at different temperatures (15 ºC, 25 ºC, and 35 ºC). P: water vapor pressure. Pₕ sat: saturation water vapor pressure at the given temperature. Water cycling stability test of 230 adsorption–desorption cycles of COF-480-hydrazide (c) subjected to an isobaric (1.7 kPa water vapor) temperature-swing adsorption between 30°C and 85 °C, corresponding to 40% and 3% RH, respectively; Full dataset reported in the SI, Section S8.
framework, given that both COFs show comparable pore diameters of around 11 Å.

It is noteworthy that even such a small conversion (9.2%) significantly shifted the water uptake to lower RH and resulted in an increase of water affinity to the framework. Thus, COF-480-hydrazide extracts water from air at even lower humidity compared to the pristine hydrazine-linked COF, AB-COF.  

To test the long-term stability of COF-480-hydrazide over multiple water adsorption-desorption cycles, we subjected the COF to an isobaric (1.7 kPa water vapor) temperature-swing adsorption between 30 °C and 85 °C, corresponding to 40% and 3% RH, respectively. The water uptake working capacity under these conditions was 0.33 g g⁻¹. After 230 adsorption-desorption cycles, we observed a reduction of the working capacity by 7% (Figure 1c). This is attributed to partial hydrolysis of the sample upon long-term exposure to water vapor at elevated temperature.

Next, we pursued the oxidation of the hydrazine-linked Py-COF-2D to COF-482-hydrazide. Noteworthy, partial oxidation of the hydrazine to the hydrazide linkage was already observed during the synthesis of the Py-COF-2D—even under inert reaction conditions (N₂ atmosphere; see SI Section S2). This phenomenon was mentioned by Lotsch and co-workers, and in our work, the observation was substantiated by FT-IR spectroscopy and ¹⁵N multi-CPMAS NMR spectroscopy on a 50% ¹⁵N-labeled sample (see SI, Sections S3 and S4). Presumably, both steric and electronic effects contribute to the more facile oxidation of Py-COF-2D as compared to AB-COF.

For further oxidation we applied comparable conditions as described for AB-COF. Similarly, we observed enhanced attenuation of the characteristic imine band at 1625 cm⁻¹ in the FT-IR spectroscopic traces and a relative increase of the carbonyl band at 1674 cm⁻¹. The solid-state ¹⁵N multi-CPMAS NMR spectroscopy on a 50% ¹⁵N-labeled sample showed an enhanced peak at 125.3 ppm, further substantiating successful oxidation. The conversion of the hydrazine- to the hydrazide-linkage amounted to 26.3% conversion (see SI, Section S4). TGA under nitrogen and air flow revealed a comparable onset in decomposition for the pristine and oxidized samples at around 500 °C.

PXRD analysis of the microcrystalline powder of Py-COF-2D and COF-482-hydrazide indicated retention of crystallinity after oxidation (see SI, section S6). Furthermore, the PXRD traces indicated a shift of the highest intensity peaks from 4.5° 2θ, associated with the (1-10) lattice plane, for the hydrazine-linked COF to 4.6° 2θ for the hydrazide COF. This observation correlates well with the anticipated changes in unit cell parameters upon oxidation (see SI, Section S6).

N₂ sorption analysis at 77 K of Py-COF-2D and COF-482-hydrazide demonstrated permanent porosity and revealed a Type I isotherm, indicating a micropore structure. The calculated BET surface areas amounted to 1850 m² g⁻¹ for the hydrazine-linked Py-COF-2D and 1700 m² g⁻¹ for COF-482-hydrazide. The reduction in specific surface area is in line with the weight increase upon oxidation. The pore size distribution, calculated from the N₂ adsorption isotherms using the above-described parameters, indicated two kinds of pores with diameters of 15.9 Å and 18.5 Å for Py-COF-2D and 15.4 Å and 18.5 Å for the oxidized COF-482-hydrazide.

Figure 2. Overlay of water isotherms of Py-COF-2D and COF-482-hydrazide at 25 °C. P: water vapor pressure. P_sat: saturation water vapor pressure.

Finally, to demonstrate postsynthetic oxidation of hydrazine- to hydrazide-linked COFs in 3D, we synthesized COF-670-hydrazine from the tetrahedral MTBA and linear hydrazine building units (Scheme 1c and Figure 3a). The structure crystallized in the dia topology with 6-fold interpenetration (see SI, Section S6). Similar to our observation with Py-COF-2D, partial oxidation of the hydrazine to hydrazide units in COF-670-hydrazine was already observed at this stage (see FT-IR and ¹⁵N spectroscopic traces, Sections S3 and S4 in the SI). This confirmed the susceptibility of the 3D COF to oxidation. Upon further oxidation, we observed enhanced

**Figure 2.** Overlay of water isotherms of Py-COF-2D and COF-482-hydrazide at 25 °C. P: water vapor pressure. P_sat: saturation water vapor pressure.
attenuation of the characteristic imine band at 1625 cm\(^{-1}\) in the FT-IR spectroscopic traces and a more intensive band at 1674 cm\(^{-1}\) (see SI, Section S3). Successful oxidation was further confirmed through \(^{15}\)N solid-state CP-MAS NMR spectroscopy on a 50% \(^{15}\)N-labeled sample before and after oxidation. The \(^{15}\)N NMR peaks at 127.8 ppm intensified upon oxidation and the conversion of the hydrazine- to the hydrazide-linkage amounted to 13.4% (Figure 3b and SI, Section S4). TGA under nitrogen and air flow revealed an increase in thermal stability upon oxidation of the hydrazine-linked COF. Here, COF-670-hydrazine and COF-670-hydrazide showed an onset in thermal decomposition at around 350 °C.

PXRD analysis of the microcrystalline powder of the hydrazine-linked COF and comparison with structural models of varying degrees of interpenetration allowed us to assign the degree of interpenetration to 6-fold (Figure 3c). After oxidation, the PXRD indicated retention of the crystallinity (Figure 3d and SI section S6). Furthermore, the PXRD traces indicated a small shift of the highest intensity peaks at 9.4° 2\(\theta\), associated with the (020) lattice plane, for the hydrazine-linked COF to 9.5° 2\(\theta\) for the hydrazide-linked COF. This observation correlated well with the anticipated changes in unit cell parameters upon oxidation (see SI, Section S6). We also observed shifts to higher 2\(\theta\) values for the designated reflections occurring between 13° to 25°.
N₂ sorption analysis at 77 K of the two samples demonstrated permanent porosity and revealed a Type I isotherm, indicating a micropore structure. The calculated BET surface areas amounted to 1209 m² g⁻¹ for the hydrazine-linked COF-670-hydrazine and 699 m² g⁻¹ for COF-670-hydrazide. The pore size distributions, calculated from the N₂ adsorption isotherms using the above-described parameters, indicated overall retention of the pore diameters of 6 Å after the oxidation (see SI, Section S7).

The pristine COF-670-hydrazine displayed a water sorption isotherm with a slightly sloped pore-filling step at around 43% RH and a total uptake of 0.33 g g⁻¹ at 25 °C (see SI, Section S8).

CONCLUSION
We synthesized a series of 2D and 3D hydrazide-linked COFs by applying a postsynthetic strategy. This involved first the synthesis of the hydrazine-linked frameworks, followed by oxidation to afford the hydrazide-linked structures, in which the synthesis and transformation from hydrazine to hydrazide was further substantiated by ¹⁵N multi-CP-MAS NMR. In addition, we subjected the frameworks to isothermal water sorption analysis. This not only allowed us to discover new structures capable of water harvesting from desert air, but also provided us with additional insight into the micro- and macroscopic changes occurring in the framework. Importantly, we found that even small modifications of the molecular structures of the frameworks can lead to impactful differences in the water isotherm profiles. Overall, this study expands the scope of COF chemistry to include crystallization of irreversible linkages and shows that these new systems can extract water from low-humidity air.

ASSOCIATED CONTENT
Supporting Information
The Supporting Information is available free of charge on the ACS Publications website.
Synthesis and full characterization of COFs including EA, FT-IR spectroscopy, NMR spectra, PXRD data, computational modeling, gas uptake measurements, TGA, and water isotherms (PDF).

Accession Codes
CCDC 2164391 contains the supplementary crystallographic data for COF-670-hydrazine. 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 Author
*Omar Yaghi: yaghi@berkeley.edu

Author Contributions
H.L.N. and C.G. conceived the idea. H.L.N, C.G., and A.M. synthesized and characterized materials. N.H. and A.L. collected and interpreted data on water vapor sorption and quantitative ¹⁵N-NMR spectra, respectively. O.M.Y directed the project. The manuscript was written through contributions of all authors. H.L.N. and C.G. contributed equally.

Funding Sources
Defense Advanced Research Projects Agency (DARPA) under Contract No: HR001-21-C-0020; Leopoldina postdoctoral fellowship of the German National Academy of Sciences (LPDS 2019-02); Studienstiftung des deutschen Volkes and Kavli ENSI Philomathia Graduate Student Fellowship.

Notes
The authors declare the following competing financial interest(s): Omar M. Yaghi is co-founder of Water Harvesting Inc., aiming at commercializing related technologies.

ACKNOWLEDGMENT
We acknowledge the financial support from the Defense Advanced Research Projects Agency (DARPA) under Contract No: HR001-21-C-0020. Any opinions, findings and conclusions or recommendations expressed in this material are those of the author(s) and do not necessarily reflect the views of the DARPA. C.G. is a Leopoldina postdoctoral fellow of the German National Academy of Science (LPDS 2019-02). N.H. thanks the Studienstiftung des deutschen Volkes and acknowledges the receipt of the Kavli ENSI Philomathia Graduate Student Fellowship. We acknowledge the College of Chemistry Nuclear Magnetic Resonance Facility for resources and staff assistance, where instruments are partially supported by NIH 5sOD024998. We thank Ms. Carlijn van Beek for synthetic contributions.

ABBREVIATIONS
COFs, covalent organic frameworks; EA, elemental analysis; FT-IR, Fourier-transform infrared spectroscopy; NMR, nuclear magnetic resonance spectroscopy; PXRD, powder X-ray diffraction; TGA, thermogravimetric analysis.

REFERENCES


