Chemical Conversion of Linkages in Covalent Organic Frameworks

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ABSTRACT: The imine linkages of two layered, porous covalent organic frameworks (COFs), TPB-TP-COF ([C₆H₃(C₆H₄N)]₂[C₆H₄(CH₂)₂], 1) and 4PE-1P-COF ([C₆(C₆H₄N)]₃[C₆H₄(CH₂)₂]₂, 2), have been transformed into amide linkages to make the respective isostuctural amide COFs 1′ and 2′ by direct oxidation with retention of crystallinity and permanent porosity. Remarkably, the oxidation of both imine COFs is complete, as assessed by FT-IR and ¹³C CP-MAS NMR spectroscopy and demonstrates (a) the first chemical conversion of a COF linkage and (b) how the usual “crystallization problem” encountered in COF chemistry can be bypassed to access COFs, such as these amides, that are typically thought to be difficult to obtain by the usual de novo methods. The amide COFs show improved chemical stability relative to their imine progenitors.

Covalent organic frameworks (COFs) are formed from molecular organic building units linked by covalent bonds to form crystalline, porous, extended solids.¹⁻³ On a fundamental level, their structure can be divided into two components: linkers (building units) and linkages (bonds formed between those units upon reticulation). The great promise of COFs is rooted in the idea that the entire structure is organic and thus may be modified through the extensive tools of molecular synthesis. Accordingly, a large number of COFs have been modified postsynthetically to introduce functionalities onto the linkers and tailor the pores.⁶⁻¹⁰ While this has expanded the applications of COFs, it is, however, the linkage and its chemistry that dictate the fundamental chemical and physical properties as well as the inherent limitations of the material. To date, direct modification of the linkage has not been achieved. Instead, the manner in which new linkages are accessed is through de novo synthesis. The intrinsic constraint this places on COF chemistry is that one is required to overcome the crystallization problem: for each new linkage, conditions must be found such that the reaction is sufficiently reversible to allow for dynamic error correction.¹¹,¹² Thus, chemistries based on linkages with limited microscopic reversibility are left unexplored.

Here we introduce an approach that bypasses the crystallization problem in making new COF linkages. Specifically, two layered imine COFs, TPB-TP-COF (1) and 4PE-1P-COF (2), were used as starting materials and subjected to mild oxidative conditions to convert the imine linkages quantitatively and give the corresponding amide-linked COFs [C₆H₃(C₆H₄NH)]₂[C₆H₄(CO₂)]₃ (1′) and [C₆(C₆H₄NH)]₃[C₆H₄(CO₂)]₂ (2′) without loss of their underlying topology, crystallinity, and permanent porosity (Figure 1). In essence, each imine framework was treated as a discrete molecule and subjected to a reaction normally carried out in molecular organic chemistry, leading to a completely new material. This method constitutes a new direction in COF chemistry where making new linkages is no longer subject to the trial and error and inherent uncertainties of de novo synthesis. The new COFs 1′ and 2′ extend the amide functionality, already prevalent and important in biological macromolecules and synthetic polymers, into the realm of porous, crystalline materials. In this context, we show that these amides exhibit chemical stability superior to that of their imine progenitors.

Supporting Information

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from 1 and 2 by treatment with sodium chlorite (11 equiv per imine functionality), acetic acid (10 equiv), and 2-methyl-2-butene (100 equiv) in dioxane for 48 h, with the sodium chlorite being added in two equal portions 24 h apart (SI, section S2).

Progression of the amidation reaction was monitored by Fourier transform infrared (FT-IR) spectroscopy (Figure 2a). The oxidized product 1’ exhibited the emergence of a C=O amide stretch (1651 cm⁻¹) and the disappearance of the C=N imine stretch (1622 cm⁻¹) found in 1. These characteristic

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**Figure 1.** (a) Conversion of imine TPB-TP-COF (1) to amide TPB-TP-COF (1’) under oxidative reaction conditions. (b) Oxidation of imine 4PE-1P-COF (2) to amide 4PE-1P-COF (2’). In the space-filling diagrams, carbon, nitrogen, oxygen, and hydrogen atoms are represented as black, blue, red, and white spheres, respectively.
imine and amide stretches were corroborated by the spectra obtained for a molecular model (SI, section S3). The FT-IR spectra of 2 and 2′ showed similar appearance and disappearance of these characteristic vibrations (Figure 2b).

Conversion of the materials from imine to amide was also observed by 13C cross-polarization magic angle spinning (CP-MAS) NMR spectroscopy. In the 13C CP-MAS NMR spectrum of 1′ at natural abundance, the imine carbon peak at 157 ppm was no longer present after oxidation. Indeed, a new peak, attributed to the amide carbonyl, was observed at 166 ppm (SI, section S4). In addition, small shifts in the frequencies of the amine and imine and amide stretches were corroborated by the spectra obtained for a molecular model (SI, section S3). The FT-IR spectra of 2 and 2′ showed similar appearance and disappearance of these characteristic vibrations (Figure 2b).

Conversion of the materials from imine to amide was also observed by 13C cross-polarization magic angle spinning (CP-MAS) NMR spectroscopy. In the 13C CP-MAS NMR spectrum of 1′ at natural abundance, the imine carbon peak at 157 ppm was no longer present after oxidation. Indeed, a new peak, attributed to the amide carbonyl, was observed at 166 ppm (SI, section S4). In addition, small shifts in the frequencies of the aromatic carbon resonances between 1 and 1′ support the oxidation of the imine functionality to the amide. Analogous results were obtained for the CP-MAS NMR spectra of 2 and 2′.

To unambiguously determine the degree of conversion from imine to amide, further 13C CP-MAS NMR experiments were performed with 13C-labeled terephthalaldehyde (50% enriched at the aldehyde carbon) to improve the signal-to-noise ratio (Figure 2c,d). This enabled monitoring of the transformation of the nitrogen-bound carbon from imine to amide for 1′ and 2′ and provided evidence for the excellent conversion of both substrates. The imine peak was absent in the 13C CP-MAS NMR spectra of the amide materials across various cross-polarization contact times between 100 μs and 7 ms.

The crystallinity of the amide material was measured by powder X-ray diffraction (PXRD). The obtained powder pattern for 1′ is in good agreement with the expected amidated structure (SI, section S5). This analysis indicates that the symmetry of 1 (space group P6, No. 168) is conserved, implying that the eclipsed stacking configuration is retained (Figure 2e). Slight changes in unit cell parameters were found for the amide (a = b = 36.54 Å, c = 4.05 Å) compared with the imine (a = b = 35.93 Å, c = 4.42 Å). The PXRD pattern of 2′ demonstrates similarly high crystallinity (Figure 2f) and agrees well with that of the predicted model structure. While the amide material provided evidence of slight unit cell changes (a = b = 38.93 Å, c = 5.45 Å compared with a = b = 38.61 Å, c = 5.82 Å for the imine), no significant structural rearrangement of the material was observed during the transformation.

Prior to structural elucidation of 1′ and 2′, all of the materials were solvent-exchanged and activated (SI, section S2). Permanent porosity of the materials was then confirmed by N2 sorption at 77 K (Figure 2g,h). Resulting Brunauer–Emmett–Teller (BET) surface areas of 1250 and 655 m2 g−1 were estimated for 1 and 1′, respectively. N2 isotherms of 2 and 2′ yielded similar results: the amidated material exhibited a lower BET surface area than the imine COF (1520 and 1190 m2 g−1 measured for the imine and amide, respectively). Part of the loss in gravimetric surface area can be attributed to an increase in framework mass and a decrease in pore volume. Additionally, 13C MAS NMR spectroscopy of 1′ imbibed with DMSO-d6 indicates the presence of included oxidized oligomers within the pores, explaining the additional surface area decrease (SI, section S4). Work is ongoing to determine the point in our synthesis at which these oligomers are generated and how best to remove them.

For both 1′ and 2′, the measurements show the type IV isotherm seen for their imine counterparts. This further confirms that the amidation procedure does not lead to significant changes in structure.

The chemical stability of the amide materials was examined by PXRD and sorption after 24 h treatment in 12 M HCl(aq) and 1 M NaOH(aq) (1 and 1′ in Figure 3; 2 and 2′ in Figure S15). The difference between the stabilities of the amine and amide materials is most striking in the case of acidic conditions, where the amides retain crystallinity while the corresponding imines are nearly or completely dissolved and the remaining material is rendered amorphous. However, a small decrease in signal-to-noise ratio and a 25–60% decrease in surface area of the amide materials indicate some structural damage upon treatment (SI, section S7). Nonetheless, the differential stability of these materials further corroborates their conversion from imines to amides.

To conclude, a method of deriving amide-linked COFs from their corresponding imine-linked frameworks has been developed. This reaction is performed under mild conditions.
and results in amide COFs retaining both crystallinity and permanent porosity. This material also exhibits enhanced stability under aqueous acidic and basic conditions relative to the corresponding imine-linked framework. This method constitutes a new direction in COF chemistry whereby the synthesis of new linkages bypasses the usual crystallization problem encountered in linking organic building blocks into extended structures by covalent bonds.

**ASSOCIATED CONTENT**

*Supporting Information*
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**Notes**
The authors declare no competing financial interest.

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**REFERENCES**


Figure 3. Comparison of diffraction patterns of (a) 1' and (b) 1 after 24 h treatment in 12 M HCl and 1 M NaOH.